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Kathleen Fuller Eric Linnell John Caive
All searchers are located in the library in CP3/4 3D62

430/2001. 2801.

1. A surface treatment agent which, when applied to a substrate prior to formation of a resist pattern thereon, strengthens adhesion between the substrate and the resist pattern, the surface treatment agent comprising at least one compound of the following compositional formula:

$$R^{1}R_{a}^{2}(OX)_{b}SiO_{(3-a-b)/2}.$$
 (1)

wherein R1 is a -(CH2), Y moiety in which Y is epoxycyclohexyl, glycidoxy,  $N-\beta$ -aminoethylamino, amino, N-phenylamino, mercapto or isocyanate, and n is an integer from 0 to 4; R2 is a monovalent hydrocarbon group of 1 to 4 carbons; X is hydrogen or a monovalent hydrocarbon group of 1 to 4 carbons; "a" is 0 or 1, and "b" is 0, 1 or 2 when "a" is 0, and "b" is 0 or 1 when "a" is 1.

2. The surface treatment agent of claim 1, wherein the 20 compound of compositional formula (1) is prepared by hydrolyzing a silane of general formula (2):

$$R^{1}R^{2}_{a}Si(OZ)_{c}$$
 (2)

- wherein R1, R2 and "a" are as defined above; Z is a 25 monovalent hydrocarbon group of 1 to 4 carbons; and "c" is a number which satisfies the condition a+c = 3.
- 3. The surface treatment agent of claim 1, wherein R1 is selected from the group consisting of  $\beta$ -(3,4-epoxy-30 cyclohexyl)ethyl, γ-aminopropyl, γ-mercaptopropyl, γisocyanatepropyl, N- $\beta$ -(aminoethyl)- $\gamma$ -aminopropyl,  $\gamma$ - $\sqrt{\text{glycidoxypropyl}}$  and N-phenyl- $\gamma$ -aminopropyl.
- A patterning process comprising the steps of applying 35 the surface treatment agent of claim 1 to a substrate and baking, then applying thereon a photoresist composition and patterning the photoresist.

10

15

- 5. The patterning process of claim 4, wherein the substrate is a metal or metal oxide substrate.
- 6. The patterning process of claim 5, wherein the metal or metal oxide making up the substrate is aluminum, iron, nickel, copper, tantalum, gold, or an oxide thereof.

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L17

L18

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(FILE 'HCAPLUS' ENTERED AT 16:23:07 ON 15 MAY 2002)
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          31585 S FUJII ?/AU
L1
L2
          66562 S KATO ?/AU
L3
            711 S L1 AND L2
L4
           7977 S FUJII T?/AU
L5
           8262 S KATO H?/AU
L6
             20 S L4 AND L5
          69162 S RESIST OR RESISTS OR PHOTORESIST?
L7
              6 S L6 AND L7
L8
                SEL L8 1-6 RN
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Ь9
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              4 S L9 AND SI/ELS
L10
                E SILICA/CN
              1 S E3
L11
L12
              3 S L10 NOT L11
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L13
             90 S L12
L14
              1 S L13 AND L7
         990221 S PATTERN? OR DESIGN OR DESIGNS OR DESIGNED OR DESIGNING#
L15
L16
              5 S L13 AND L15
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10450 S (SUBSTRAT? OR SURFACE? OR BASE# OR SUBSTRUCT? OR UNDERS

16425 S (TREAT? OR PRETREAT? OR CONDITION? OR PRECONDITION? OR

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L19 172333 S L18(2A)L17
            7 S L13 AND L19
L20
   FILE 'REGISTRY' ENTERED AT 16:40:58 ON 15 MAY 2002
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              _____
L21
              STR
             _____
L22
           50 S L21
        6214 S L21 FUL
L23
               DEL SAG001/Q
               SAV TEM L23 SAG001/A
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          609 S L28 NOT N/ELS
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L31 3011 S L30
L32
         8543 S L29
L33
         9057 S L28
        21288 S L23
L34
           27 S L31 AND L7
L35
L36
            9 S L35 AND (L15 OR L19)
          113 S L32 AND L7
L37
L38
           54 S L37 AND (L15 OR L19)
            7 S L37 AND L15 AND L19
7 S L33 AND L7 AND L15 AND L19
L39
L40
           15 S L34 AND L7 AND L15 AND L19
L41
           23 S L14 OR L16 OR L20 OR L36 OR L39 OR L40
L42
            8 S L41 NOT L42
L43
    FILE 'REGISTRY' ENTERED AT 18:05:09 ON 15 MAY 2002
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=> d 123 que stat L21 S'

STR

6214 ANSWERS

## Page 1-A

034 N→ Ph S N= C= O 030 31 E1 037 38 39

Page 1-B
REP G1=(0-4) CH2
VAR G2=8/12/18/NH2/30/34/37
VAR G3=0/25
NODE ATTRIBUTES:
HCOUNT IS E1 AT 34
NSPEC IS RC AT 25
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 26

STEREO ATTRIBUTES: NONE

L23 6214 SEA FILE=REGISTRY SSS FUL L21

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=> d 142 1-23 cbib abs hitstr hitind

L42 ANSWER 1 OF 23 HCAPLUS COPYRIGHT 2002 ACS
2002:169372 Document No. 136:220632 Method of forming metal film
pattern on glass substrate. Hayashi, Shigehiko; Iwamoto,
Masahiro; Kuroda, Kotaro; Igarashi, Tetsuya (Mitsuboshi Belting
Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2002068782 A2 20020308, 7
pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2000-258425
20000829.

The process comprises the steps of (1) covering an area where a pattern will not be formed with a resist film, (2) surface\_treating with a silane coupling agent having a functional group selected from mercapto, amino, cyano, amido, and ureido, (3) fixing metal particles on the glass substrate via the silane coupling agent on the surface, (4) electroless-plating with a metal, and (5) removing the resist pattern

IT 4420-74-0, 3-Mercaptopropyltrimethoxysilane (method of forming metal film pattern on glass substrate)

RN 4420-74-0 HCAPLUS

CN 1-Propanethiol, 3-(trimethoxysilyl)- (7CI, 8CI, 9CI) (CA INDEX NAME)

IC ICM C03C017-38 ICS C23C018-18; C23C018-31

CC 57-1 (Ceramics)
Section cross-reference(s): 42

IT Coating process

(electroless; method of forming metal film pattern on
glass substrate)

IT 7440-57-5, Gold, processes

(method of forming metal film **pattern** on glass substrate)

IT 60676-86-0, Quartz glass

(method of forming metal film pattern on glass

substrate)

- 1T 4420-74-0, 3-Mercaptopropyltrimethoxysilane 7440-02-0, Nickel, uses 138988-79-1, Microposit XP 90104A 273204-64-1, Microposit S 1808 (method of forming metal film pattern on glass substrate)
- L42 ANSWER 2 OF 23 HCAPLUS COPYRIGHT 2002 ACS
- 2002:169356 Document No. 136:220624 Manufacture of optical devices with convex lenses on glass substrates by sol-gel process using mask patterns. Yokoyama, Masaaki; Sakurai, Yoshiaki; Nagayama, Tomoo (Japan). Jpn. Kokai Tokkyo Koho JP 2002068755 A2 20020308, 7 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2000-265163 20000901.
- The process involves (A) forming layers of organosilicon resins, preferably polysilanes, on glass substrates, (B) irradiating radiation on the layers selectively on regions where convex lenses are to be formed, (C) bringing the layers into contact with liq. gels of the convex lens-forming glass compns. to deposit the gels selectively on the irradiated regions, and (D) drying the deposited liq. gels and heating the obtained dry gels for vitrification. Between the steps of C and B, the liq. gel-deposited glass substrates may be brought into contact with solns. contg. metals, org. acids, or inorg. acids or water to control refractive indexes of the resulting lenses. The radiation may be electron beam, x-ray, or UV. The liq.gel may be prepd. from .gtoreq.1 metal alkoxides. The optical devices may be microlens arrays for charge coupled devices (CCD), etc.

29159-37-3P, 3-Aminopropyltriethoxysilane homopolymer (manuf. of microlenses by sol-gel process using mask patterns of organosilicon resins)

RN 29159-37-3 HCAPLUS

CN 1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 919-30-2

CMF C9 H23 N O3 Si

```
OEt.
EtO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>
     OEt
IC
     ICM
         C03B008-02
     ICS
          G02B003-00
     57-1 (Ceramics)
CC
     Section cross-reference(s): 73
ST
     optical lense convex sol gel lithog; microlens array manuf sol gel
     lithog; polysilane pattern sol gel microlens device;
     silica qel polysilane pattern microlens array
IT
     Polysilanes
        (fluorine-contg.; manuf. of microlenses by sol-gel process using
        mask patterns of)
IT
     Microlenses
     Photolithography
     Sol-gel processing
        (manuf. of microlenses by sol-qel process using mask
        patterns)
IT
     Polysilanes
        (manuf. of microlenses by sol-qel process using mask
        patterns of)
     Fluoropolymers, uses
IT
        (polysilane-; manuf. of microlenses by sol-gel process using mask
        patterns of)
TT
     31324-77-3
                  402496-26-8
        (manuf. of microlenses by sol-qel process using mask
        patterns of)
     11099-06-2P, Tetraethoxysilane homopolymer 29159-37-3P,
IT
     3-Aminopropyltriethoxysilane homopolymer
        (manuf. of microlenses by sol-gel process using mask
        patterns of organosilicon resins)
     ANSWER 3 OF 23 HCAPLUS COPYRIGHT 2002 ACS
             Document No. 136:126537 Method for pattern
     formation using organic self-assembled monolayer and UV light and
     method for forming organic self-assembled monolayers therefor.
     Ishida, Masaya (Seiko Epson Corp., Japan). Jpn. Kokai Tokkyo Koho
     JP 2002019008 A2 20020122, 5 pp. (Japanese). CODEN: JKXXAF.
     APPLICATION: JP 2000-202200 20000704.
     The title method includes the steps of: forming a first org.
AB
     self-assembled monolayer of .ltoreq.3 nm thickness on a substrate;
     and forming a second org. self-assembled monolayer having different
     functional groups from the first org. self-assembled monolayer.
     method, which has the two monolayers instead of one monolayer,
     provides the photomask which shortening the exposure time for
     patterning
     14814-09-6
IT
```

```
(monolayer material for forming org. self-assembled monolayers)
     14814-09-6 HCAPLUS
RN
CN
     1-Propanethiol, 3-(triethoxysilyl)- (7CI, 8CI, 9CI) (CA INDEX NAME)
     OEt
EtO-Si-(CH_2)_3-SH
     OEt
IC
     ICM
          B32B009-00
          B05D001-36; B05D003-06; G03F007-004; G03F007-20; G03F007-36;
     ICS
          H01L021-027
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and
CC
     Other Reprographic Processes)
     patterning org self assembled monolayer
ST
     Light-sensitive materials
IT
       Řesists
     Self-assembled monolayers
        (method for patterning using org. self-assembled
        monolayer and method for forming org. self-assembled monolayer)
     14814-09-6
                  101947-16-4, Heptadecafluoro-1,1,2,2-
IT
     tetrahydrodecyltriethoxysilane
        (monolayer material for forming org. self-assembled monolayers)
     ANSWER 4 OF 23 HCAPLUS COPYRIGHT 2002 ACS
L42
              Document No. 136:12840 Photosensitive composition
2001:864938
     containing adhesive aid. Kasumi, Kenichi; Yumiba, Tomoyuki;
     Fujimoto, Koji (Toray Industries, Inc., Japan). Jpn. Kokai Tokkyo
     Koho JP 2001330962 A2 20011130, 10 pp. (Japanese). CODEN: JKXXAF.
     APPLICATION: JP 2000-146224 20000518.
     The compn., comprising a polymer, an adhesive aid with metals, and a
AB
     photosensitive agent, is coated on a metal foil, pre-baked,
     imagewise exposed to form patterns and has av. heat linear
     expansion coeff. .ltoreq.25 .times. 10-6(/.degree., at
     30-100.degree.) after curing.
                                    The compn. shows good adhesion with
     metals and clear patterns are formed.
     4420-74-0
IT
        (photosensitive compn. contg. adhesive aid)
```

1-Propanethiol, 3-(trimethoxysilyl)- (7CI, 8CI, 9CI)

OMe 
$$\mid$$
 MeO-Si-(CH<sub>2</sub>)<sub>3</sub>-SH  $\mid$  OMe

NAME)

RN

CN

4420-74-0 HCAPLUS

```
IC
     ICM
         G03F007-085
          C08K005-34; C08K005-36; C08L101-00; G03F007-037; G03F007-038;
     ICS
          G03F007-40; H01L021-027; H01L021-312
CC
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and
     Other Reprographic Processes)
     Section cross-reference(s): 38
     photoresist adhesive aid metal surface; linear expansion
ST
     coeff photoresist polyamide
     Photoresists
IT
        (photosensitive compn. contg. adhesive aid)
     66-71-7, 1,10-Phenanthroline 95-14-7, 1,2,3-Benzotriazole
IT
     120-78-5, Dibenzothiazyl disulfide
                                          148-24-3, 8-Quinolinol, uses
     149-30-4, 2-Mercaptobenzothiazole
4420-74-0 13733-91-0, 2-Anilino-4,
                                          288-94-8, 1H-Tetrazole
                 13733-91-0, 2-Anilino-4,6-dimercapto-s-triazine
        (photosensitive compn. contq. adhesive aid)
     ANSWER 5 OF 23 HCAPLUS COPYRIGHT 2002 ACS
L42
              Document No. 135:160159 Surface
2001:582290
     treatment agent for resist pattern, and
     patterning process. Fujii, Toshihiko; Kato, Hideto (Japan).
       U.S. Pat. Appl. Publ. US 20010012601 A1 20010809, 4 pp.
     (English). CODEN: USXXCO. APPLICATION: US 2001-772001 20010130.
     PRIORITY: JP 2000-21103 20000131.
     A surface treatment agent comprises an
AB
     oligomeric or polymeric silicon-contg. compd. having an
     epoxycyclohexyl, glycidoxy, N-.beta.-aminoethylamino, amino,
     N-phenylamino, mercapto or isocyanate group. When the
     surface treatment agent is applied to a metal or
     metal oxide substrate prior to formation of a resist
     pattern thereon, it serves to strengthen adhesion between
     the substrate and the photoresist film.
                 .gamma.-Aminopropyltriethoxysilane polymer
     29159-37-3P
IT
     191093-22-8P, 3-Aminopropyltriethoxysilane-3-
     Mercaptopropyltriethoxysilane copolymer 352668-65-6P
        (surface treatment agent; Si-contq. compd.
        surface treatment agent for resist
        patterning)
RN
     29159-37-3
                HCAPLUS
     1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX
CN
     NAME)
     CM
          1
     CRN
          919-30-2
     CMF
          C9 H23 N O3 Si
```

14814-09-6

C9 H22 O3 S Si

CRN CMF

```
OEt
EtO-Si-(CH<sub>2</sub>)<sub>3</sub>-SH
     OEt
IC
     ICM G03C005-00
NCL
     430322000
CC
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and
     Other Reprographic Processes)
     Section cross-reference(s): 35, 38, 76
     photoresist patterning silicon oligomeric
ST
     adhesion surface treatment agent
     Photolithography
IT
       Photoresists
        (Si-contg. compd. surface treatment agent for
        resist patterning)
     7429-90-5, Aluminum, uses
                                 7440-50-8, Copper, uses
IT
        (substrate; Si-contg. compd. surface treatment
        agent for resist patterning)
     29159-37-3P, .gamma.-Aminopropyltriethoxysilane polymer 191093-22-8P, 3-Aminopropyltriethoxysilane-3-
TT
     Mercaptopropyltriethoxysilane copolymer 352668-65-6P
        (surface treatment agent; Si-contg. compd.
        surface treatment agent for resist
        patterning)
L42
     ANSWER 6 OF 23 HCAPLUS COPYRIGHT 2002 ACS
2001:354321 Document No. 135:93330 On the properties of an ion
     conductive system incorporated in hybrid films. Gonzalez, G.;
     Retuert, P. J.; Fuentes, S. (Departments of Chemistry, Faculty of
     Science, Universidad de Chile, Santiago, Chile). Materials Research
     Society Symposium Proceedings, 628 (Organic/Inorganic Hybrid
     Materials), CC6.19.1-CC6.19.6 (English) 2001. CODEN: MRSPDH.
     0272-9172.
                 Publisher: Materials Research Society.
     Blending the biopolymer chitosan (CHI) with
AB
     poly(aminopropilsiloxane) oligomers (pAPS), and poly(ethylene oxide)
     (PEO) in the presence of lithium perchlorate lead to ion conducting
     products whose cond. depends on the compn. of the mixt. A ternary
     phase diagram for mixts. contg. 0.2 M LiClO4 shows a zone in which
     the phys. properties of the products - transparent, flexible, mech.
     robust films - indicate a high degree of mol. compatibilization of
     the components. Comparison of these films with binary CHI-pAPS
     nanocomposites as well as the microscopic aspect, thermal behavior,
     and X-ray diffraction pattern of the product with the
     compn. PEO/CHI/pAPS/LiClO4 1:0.5:0.6:0.2 molar ratio indicates that
     these films may be described as a layered nanocomposite.
                                                                 In this
```

composite, lithium species coordinated by PEO and pAPS should be inserted into chitosan layers. Electrochem. impedance spectroscopy measurements indicate the films are pure ionic conductors with a

maximal bulk cond. of 1.7.times.10-5 Scm-1 at 40.degree.C and a sample-electrode interface capacitance of about 1.2.times.10-9 F. 29159-37-3, 3-Aminopropyltriethoxysilane homopolymer IT (properties of poly(ethylene oxide)-chitosan-aminopropyl silsesquioxane-lithium perchlorate ionic conductive system) 29159-37-3 HCAPLUS RN1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX CN NAME) CM 1 919-30-2 CRN CMF C9 H23 N O3 Si OEt Eto-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>OEt 37-5 (Plastics Manufacture and Processing) CC Section cross-reference(s): 44 7439-93-2D, Lithium, complexes with poly(ethylene oxide), properties IT 9012-76-4, Chitosan 25322-68-3D, Poly(ethylene oxide), lithium complexes 29159-37-3, 3-Aminopropyltriethoxysilane homopolymer 161376-90-5, 3-Aminopropyltriethoxysilane homopolymer ladder sru (properties of poly(ethylene oxide)-chitosan-aminopropyl silsesquioxane-lithium perchlorate ionic conductive system) ANSWER 7 OF 23 HCAPLUS COPYRIGHT 2002 ACS Document No. 133:44462 Relationship between Composition 2000:386597 and Structure in Chitosan-based Hybrid Films. Fuentes, S.; Retuert, P. J.; Ubilla, A.; Fernandez, J.; Gonzalez, G. (Department of Chemistry Faculty of Science, University of Chile, Santiago, Chile). Biomacromolecules, 1(2), 239-243 (English) 2000. CODEN: BOMAF6. ISSN: 1525-7797. Publisher: American Chemical Society. Chitosan/poly(aminopropylsiloxane) hybrid films were obtained by AB blending 3-(aminopropyl)siloxane oligomers (pAPS) with chitosan The pAPS oligomers were prepd. by the sol-gel method starting from 3-(aminopropyl)triethoxysilane. These hybrids were characterized by chem., spectroscopic and morphol. methods. Scanning electron micrographs of hybrid films of different compn. revealed an organized microscopic pattern suggesting the existence of systematic interactions among their components. Comparison of the thermal stabilities and X-ray diffraction patterns as well as FT-IR spectra of the films with those of the pure components revealed that nanocomposites were formed.

Similar studies of films including lithium perchlorate, as a third

component, showed that addn. of certain amt. of lithium ions

affected the structure of the CHI/pAPS films. When addn. of the lithium salt exceeded the homogeneous incorporation limit, a little excess generated anisotropically oriented **patterns** in the hybrid films.

29159-37-3, 3-(Aminopropyl)triethoxysilane homopolymer (relationship between compn. and structure in chitosan-based hybrid films)

RN 29159-37-3 HCAPLUS

CN 1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 919-30-2

CMF C9 H23 N O3 Si

CC 38-3 (Plastics Fabrication and Uses)
Section cross-reference(s): 37, 44

IT 9012-76-4, Chitosan 29159-37-3, 3(Aminopropyl)triethoxysilane homopolymer
(relationship between compn. and structure in chitosan-based hybrid films)

L42 ANSWER 8 OF 23 HCAPLUS COPYRIGHT 2002 ACS
1998:498147 Document No. 129:217996 Surface modification methods for
water and oil repellency. Kai, Yoshiaki (Matsushita Electric
Industrial Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 10204421 A2
19980804 Heisei, 6 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP

1997-12193 19970127.

Substrates are treated with F-free silane coupling agents and F-contg. carboxylic acid halides. Thus, soda glass was treated with bis(2-hydroxyethyl)-3-aminopropyltriethoxysilane hydrolyzate and acid chloride of Krytox 157FSH.

29159-37-3DP, Poly(3-aminopropyltriethoxysilane), reaction products with perfluorohexyldecanoic acid chloride (water- and oil-repellent coatings on glass and ceramics and metals and plastics)

RN 29159-37-3 HCAPLUS

CN 1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 919-30-2 CMF C9 H23 N O3 Si

IC ICM C09K003-18

C03C017-32; C04B041-83

42-10 (Coatings, Inks, and Related Products) CC Section cross-reference(s): 55, 56, 57

29159-37-3DP, Poly(3-aminopropyltriethoxysilane), reaction IT products with perfluorohexyldecanoic acid chloride 29226-47-9DP, reaction products with perfluorostearic acid chloride 90168-44-8DP, reaction products with perfluoropolyoxyalkylene carboxylic acid chloride 146221-69-4DP, reaction products with bis(hydroxyethyl)aminopropyltriethoxysilane hydrolyzate 212139-97-4DP, reaction products with aminopropyltriethoxysilane 212140-00-6DP, reaction products with (aminoethyl) aminopropyltrimethoxysilane hydrolyzates 212140-02-8DP, reaction products with perfluoro(methyldimethylbutyl) methyldimethyloctanoic acid chloride 212140-03-9DP, reaction products with aminophenyltrimethoxysilane hydrolyzates (water- and oil-repellent coatings on glass and ceramics and metals and plastics)

ANSWER 9 OF 23 HCAPLUS COPYRIGHT 2002 ACS

Document No. 126:164265 Photosolder resist 1997:154648 composition containing photocurable resin and silicone prepolymer. Kubota, Hiroyuki (Toyo Ink Mfg Co, Japan). Jpn. Kokai Tokkyo Koho JP 08320564 A2 19961203 Heisei, 9 PP. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1995-126174 19950525.

The compn. contains (A) a resin contg. CO2H and alc. OH which is AB cured by irradn. with active rays and (B) a silicone prepolymer. (B) is preferably a condensate and/or partial condensate having .gtoreq.1 functional group selected from vinyl, amino, mercapto, epoxy, and (meth)acryloyl groups. (A) preferably contains (A1) adducts of (a) an unsatd. monocarboxylic acid polymer or its copolymer with other ethylenic compd. with (b) a compd. having an epoxy group and .gtoreq.1 ethylenic bond and (A2) half esters of (c) a homopolymer of an unsatd. polybasic acid or its anhydride their copolymer with other ethylenic compd. with (d) a monohydric alc. having ethylenic bond or a polyhydric alc. having no ethylenic bond. The resist compn., useful for manuf. of printed circuit boards, provides insulating patterns with good resistance to heat, chems., water, moisture, and solvents, and. (A) is previously treated with (B) in part. 168269-79-2P

IT

(photosolder resist compn. contq. photocurable resin and silicone prepolymer) 168269-79-2 HCAPLUS RN 1-Propanethiol, 3-(trimethoxysilyl)-, polymer with CNtrimethoxymethylsilane (9CI) (CA INDEX NAME) CM CRN 4420-74-0 CMF C6 H16 O3 S Si OMe MeO-Si-(CH<sub>2</sub>)<sub>3</sub>-SHOMe CM2 CRN 1185-55-3 CMF C4 H12 O3 Si OMe MeO-Si-Me OMe 186546-51-0P IT (photosolder resist compn. contq. photocurable resin and silicone prepolymer) 186546-51-0 HCAPLUS RN 2-Propenoic acid, 2-methyl-, methyl ester, polymer with CN oxiranylmethyl 2-methyl-2-propenoate, 2-propenoic acid, trimethoxymethylsilane and 3-(trimethoxysilyl)-1-propanethiol (9CI)

CM 1

CRN 4420-74-0 CMF C6 H16 O3 S Si

(CA INDEX NAME)

$$\begin{array}{c} \text{OMe} \\ | \\ \text{MeO-Si-} (\text{CH}_2)_3 - \text{SH} \\ | \\ \text{OMe} \end{array}$$

CM 2

CRN 1185-55-3 CMF C4 H12 O3 Si

CM 3

CRN 106-91-2 CMF C7 H10 O3

CM 4

CRN 80-62-6 CMF C5 H8 O2

$$\begin{array}{ccc} ^{H_2C} & \text{O} \\ \parallel & \parallel \\ \text{Me-} & \text{C-} & \text{C-} & \text{OMe} \end{array}$$

CM 5

CRN 79-10-7 CMF C3 H4 O2

```
0
HO-C-CH-CH2
IC
     ICM
          G03F007-075
     ICS
          G03F007-038; H05K003-06
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and
CC
     Other Reprographic Processes)
     solder photoresist photocurable resin; silicone prepolymer
ST
     solder photoresist; photosolder resist
     photocurable resin
IT
     Silsesquioxanes
        (Me Ph; photosolder resist compn. contq. photocurable
        resin and silicone prepolymer)
     Solder resists
IT
        (photoresists; photosolder resist compn.
        contg. photocurable resin and silicone prepolymer)
IT
     Silsesquioxanes
        (photosolder resist compn. contq. photocurable resin
        and silicone prepolymer)
     Photoresists
IT
        (solder; photosolder resist compn. contg. photocurable
        resin and silicone prepolymer)
                   162816-07-1P, Methyltrimethoxysilane-
IT
     phenyltrimethoxysilane copolymer 168269-79-2P
        (photosolder resist compn. contq. photocurable resin
        and silicone prepolymer)
                    186546-50-9P 186546-51-0P
     156637-69-3P
                                                  186546-53-2P
IT
     186708-37-2P
        (photosolder resist compn. contq. photocurable resin
        and silicone prepolymer)
     9005-12-3, Poly[oxy(methylphenylsilylene)]
IT
        (photosolder resist compn. contq. photocurable resin
        and silicone prepolymer)
     ANSWER 10 OF 23 HCAPLUS COPYRIGHT 2002 ACS
              Document No. 126:97499 Fabrication and packaging of mesa
1996:642823
              van Hal, R. E. G.; Bergveld, P.; Engbersen, J. F. J. (MESA
     Research Institute, University Twente, Enschede, 7500, Neth.).
     Sens. Mater., 8(7), 455-468 (English) 1996.
                                                   CODEN: SENMER.
     0914-4935. Publisher: Scientific Publishing Division of MYU K.K..
     A novel three-dimensional structure for the ISFET is developed,
AB
     which simplifies the encapsulation procedure and enables flat
     packaging. The chem. and elec. characteristics of the ISFET are
     unchanged by the new structure. The ISFET is encapsulated in a
     self-aligning process with a Kapton foil by polymer bonding, while
     at the same time, the source and drain pads are connected to the
     preprocessed copper leads on the Kapton foil. 29159-37-3D, 3-Aminopropyl triethoxysilane homopolymer,
IT
```

oligomerized (silica surface treatment agent; fabrication and packaging of mesa ISFETs) RN 29159-37-3 HCAPLUS CN 1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX CM CRN 919-30-2 C9 H23 N O3 Si CMF OEt Eto-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>OEt CC 76-3 (Electric Phenomena) 29159-37-3D, 3-Aminopropyl triethoxysilane homopolymer, IT oligomerized (silica surface treatment agent; fabrication and packaging of mesa ISFETs) ANSWER 11 OF 23 HCAPLUS COPYRIGHT 2002 ACS Document No. 123:145576 Surface modification of glass 1995:744365 fibers by silane coupling agents and its effect on the strength of the interfacial boundary and properties of polypropylene glass-reinforced plastics. Zelentskii, A. N.; Gorbatkina, Yu. A.; Kuperman, A. M.; Pirogov, O. N.; Tovmasyan, M. A.; Denisov, K. A.; Vasil'ev, I. A. (Semenov Inst. Chem. Physics, Russian Acad. Sci., Moscow, 117977, Russia). Vysokomol. Soedin., Ser. A Ser. B, 37(5), 775-80 (Russian) 1995. CODEN: VSSBEE. Ozonometry was used for quant. anal. of functional groups forming on AB the surface of glass fibers as a result of the condensation polymn. of silane coupling agents. A setup was built and technique for ozonometric measurements was developed. The method was used to investigate the surface of glass fibers modified by vinyl- and aminosilanes. Anal. showed that the best results are achieved by treating the surface with vinyltriethoxysilane. In this case, coupling between the glass fibers and the polypropylene matrix and the toughness of polypropylene composites increase. 29159-37-3, .gamma.-Aminopropyltriethoxysilane homopolymer IT (surface modification of glass fibers by silane coupling agents and its effect on properties of reinforced polypropylene) 29159-37-3 HCAPLUS RN 1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX CN NAME)

CM 1

CRN 919-30-2 CMF C9 H23 N O3 Si

OEt 
$$|$$
 EtO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>  $|$  OEt

CC 37-6 (Plastics Manufacture and Processing) Section cross-reference(s): 38

L42 ANSWER 12 OF 23 HCAPLUS COPYRIGHT 2002 ACS

1995:705333 Document No. 123:97953 Disulfide-containing organic silane-modified surface for photo-patterning. Ichinose, Nobuyuki (Shingijutsu Kaihatsu Jigyodan, Japan). Jpn. Kokai Tokkyo Koho JP 07084371 A2 19950331 Heisei, 4 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1993-232082 19930917.

AB Disulfide org. group-substituted silanes are adsorbed on a surface via O bonds and the disulfide org. groups are capable of cleaving upon photoirradn. The above silanes are prepd. by reacting mercapto group substituted silane derivs. with sulfenic acid halide or disulfide compd.

IT 4420-74-0, 3-Mercaptopropyltrimethoxysilane (monolayer film from)

RN 4420-74-0 HCAPLUS

CN 1-Propanethiol, 3-(trimethoxysilyl)- (7CI, 8CI, 9CI) (CA INDEX NAME)

OMe 
$$\mid$$
 MeO-Si-(CH<sub>2</sub>)<sub>3</sub>-SH  $\mid$  OMe

IC ICM G03F007-075 ICS G03F007-26

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 76

ST monolayer film photoresist disulfide silane

IT Molecular electronic devices

(disulfide-contg. org. silane-modified oxidized surface for photo-patterning)

IT Resists

(photo-, disulfide-contg. org. silane-modified oxidized surface for photo-patterning)

IT 882-33-7, Diphenyldisulfide 4420-74-0,

3-Mercaptopropyltrimethoxysilane
 (monolayer film from)

L42 ANSWER 13 OF 23 HCAPLUS COPYRIGHT 2002 ACS

1995:604470 Document No. 123:22405 Oxides modified with organosilane compounds having acylalkyl sulfide linkage, their preparation, and photopatterned surface of the oxides. Ichinose, Nobuyuki (Shingijutsu Kaihatsu Jigyodan, Japan). Jpn. Kokai Tokkyo Koho JP 07084372 A2 19950331 Heisei, 3 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1993-232055 19930917.

The oxides have a monomol. layer of silane substituted with org. AB group having acylalkyl sulfide linkage. The modified oxides are prepd. by coating of surface of solid oxides with a monomol. layer of silane substituted with SH group followed by acylalkylation. Patterned oxides, where the acylalkyl groups are cleaved by **Patterned** patternwise irradn., is also claimed. oxide, where amino polymer mols. are bound to the unirradiated part, is also claimed. The organosilane monolayer can be patterned by generally-used near-UV light irradn. app. and the surface further modified with peptides and aminopolymers, e.g. proteins, through the organosilane are useful for ultra-thin-layer photoresists, mol. devices, biosensors, and biocompatible A quartz substrate was treated with materials. (MeO) 3Si(CH2) 3SH followed by ICH2CO2H or maleic anhydride to give modified surface, which was patternwise irradiated with KrF excimer laser and further treated with poly(ethyleneimine) to give its pattern.

IT 4420-74-0, (3-Mercaptopropyl)trimethoxysilane (quartz coating with; photopatterning of oxide surface coated with organosilane having acylalkyl sulfide linkage and modification of the surface with aminopolymers for electronic

RN 4420-74-0 HCAPLUS

CN 1-Propanethiol, 3-(trimethoxysilyl)- (7CI, 8CI, 9CI) (CA INDEX NAME)

devices and biocompatible materials)

IC ICM G03F007-075 ICS G03F007-075

- CC 74-13 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
  Section cross-reference(s): 29, 66
- ST oxide photopatterning silane coupling agent; acylalkyl sulfide linkage coupling agent; aminopolymer patternwise fixation coupling agent; protein patternwise immobilization coupling agent
- IT Oxides, processes

(substrates; photopatterning of oxide surface coated with organosilane having acylalkyl sulfide linkage and modification of the surface with aminopolymers for electronic devices and biocompatible materials)

- 1T 4420-74-0, (3-Mercaptopropyl)trimethoxysilane (quartz coating with; photopatterning of oxide surface coated with organosilane having acylalkyl sulfide linkage and modification of the surface with aminopolymers for electronic devices and biocompatible materials)
- 7631-86-9, Silica, processes
  (substrate; photopatterning of oxide surface coated
  with organosilane having acylalkyl sulfide linkage and
  modification of the surface with aminopolymers for electronic
  devices and biocompatible materials)
- L42 ANSWER 14 OF 23 HCAPLUS COPYRIGHT 2002 ACS

  1994:278208 Document No. 120:278208 Electroless process for the formation of metal **patterns** on glass. Van der Putten,

  Andreas M. T. P.; Willard, Nicolaas Petrus; Fokkink, Lambertus

Gerrit Jan; Camps, Ivo Godfried Jozef (Philips Electronics N.V., Neth.). Eur. Pat. Appl. EP 577187 A1 19940105, 8 pp. DESIGNATED STATES: R: DE, FR, GB. (English). CODEN: EPXXDW. APPLICATION: EP 1993-201804 19930622. PRIORITY: EP 1992-201925 19920629.

The process comprises modifying the substrates with a silane layer, locally removing the layer with a laser or UV-03 treatment, selectively nucleating the remaining silane layer with a polymer-stabilized Pd sol, and metalizing the substrates by electroless process. This process employs neither photoresists nor org. solvents. The method is esp. suitable for forming a black matrix on passive plates for LCD's, and on other flat, color display panels, e.g., flat cathode-ray tubes.

IT 1760-24-3P

(modification with, of glass plates, in metal **pattern** formation by electroless coating process)

- RN 1760-24-3 HCAPLUS
- CN 1,2-Ethanediamine, N-[3-(trimethoxysilyl)propyl]- (9CI) (CA INDEX NAME)

```
OMe
MeO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH-CH<sub>2</sub>-CH<sub>2</sub>-NH<sub>2</sub>
     OMe
IC
     ICM
         C03C017-38
     ICS
          G03F007-075
     57-1 (Ceramics)
CC
     Section cross-reference(s): 75
     metal pattern formation glass oxide; silane coating
ST
     removal laser pattern; polymer stabilizer palladium sol
     silane; electroless metalization glass
IT
    Silanes
        (alkoxyamino, modification with, of glass plates, in metal
        pattern formation by electroless coating process)
     Polymers, uses
IT
        (palladium sols stabilized with, activation with, of
        patterned UV-ozone-treated silane-coated glass plates,
        for metal pattern formation by electroless coating
        process)
IT
     Ultraviolet radiation
        (removal by, in presence of ozone, of silanes, in metal
        pattern formation on glass plates by electroless coating
        process)
IT
     Cathode-ray tubes
        (color, black metal matrix formation on glass plates for,
        activated pattern formation in)
IT
     Coating process
        (electroless, metalization, of glass plates, activated
        pattern formation for, for liq.-crystal display devices
        and flat color cathode-ray tubes)
IT
     Optical imaging devices
        (lig.-crystal, black metal matrix formation on glass plates for,
        activated pattern formation in)
IT
     Glass, oxide
        (plate, borosilicate, metal pattern formation on,
        electroless process for, coating with silane and
        patterned UV-ozone radiation and activation with
        palladium sol in)
     7440-02-0, Nickel, uses
IT
        (coating with, electroless, of glass plates, activated
        pattern formation for, for liq.-crystal display devices
        and flat color cathode-ray tubes)
     7440-05-3P, Palladium, properties
IT
        (colloidal, activation with polymer-stabilized, of
        patterned UV-ozone-treated silane-coated glass plates,
        for metal pattern formation by electroless coating
        process)
     919-30-2P, 3-Aminopropyl triethoxysilane 1760-24-3P
IT
```

(modification with, of glass plates, in metal **pattern** formation by electroless coating process)

- IT 9002-89-5P, Poly(vinyl alc.) 9003-39-8P, Poly(vinyl pyrrolidone)
  (palladium sols stabilized with, activation with, of

  patterned UV-ozone-treated silane-coated glass plates,
  for metal pattern formation by electroless coating
  process)
- L42 ANSWER 15 OF 23 HCAPLUS COPYRIGHT 2002 ACS
  1994:231677 Document No. 120:231677 Top-surface imaging using
  selective electroless metalization of patterned monolayer
  films. Calvert, J. M.; Dressick, W. J.; Dulcey, C. S.; Chen, M. S.;
  Goerger, J. H.; Stenger, D. A.; Koloski, T. S.; Calabrese, G. S.
  (Nav. Res. Lab., Washington, DC, 20375-5320, USA). ACS Symp. Ser.,
  537(Polymer for Microelectronics), 210-19 (English) 1994. CODEN:
- ACSMC8. ISSN: 0097-6156.

  AB A top surface imaging microlithog. process that involves selective electroless (EL) metalization of surfaces modified with ligating organosilane ultrathin films (UTFs) is described. Fabrication of metal features with 0.4 .mu.m linewidths using 193 nm exposure is shown. Metal-ligand complexation chem. is used for covalent attachment of a Pd(II) catalyst to the UTF-treated surface. The mol. nature of the UTF layer is shown to control the adhesive strength of the EL metal deposit; values of >500 psi on single crystal Si wafers have been obtained. The ligand-based UTF process is a promising approach for a range of microelectronic applications where high resoln., adherent, selective metalization is required.

1760-24-3, N-(2-Aminoethyl)-3-aminopropyltrimethoxysilane 35141-30-1 84127-79-7

(electroless metalization of surfaces modified by, in photolithog. top-surface imaging)

RN 1760-24-3 HCAPLUS

CN 1,2-Ethanediamine, N-[3-(trimethoxysilyl)propyl]- (9CI) (CA INDEX NAME)

OMe 
$$| \\ \text{MeO-Si-} (\text{CH}_2)_3 - \text{NH-CH}_2 - \text{CH}_2 - \text{NH}_2 \\ | \\ \text{OMe} \\$$

RN 35141-30-1 HCAPLUS

CN 1,2-Ethanediamine, N-(2-aminoethyl)-N'-[3-(trimethoxysilyl)propyl]-(9CI) (CA INDEX NAME)

OMe 
$$| \\ \text{MeO-Si-} (\text{CH}_2)_3 - \text{NH-CH}_2 - \text{CH}_2 - \text{NH-CH}_2 - \text{CH}_2 - \text{NH}_2 \\ | \\ \text{OMe} \\$$

RN 84127-79-7 HCAPLUS

CN 2-0xa-7,10-diaza-3-siladodecan-12-oic acid, 7,10-bis(carboxymethyl)-3,3-dimethoxy- (9CI) (CA INDEX NAME)

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

IT Resists

(photo-, ligating organosilanes, selective electroless metalization of)

1760-24-3, N-(2-Aminoethyl)-3-aminopropyltrimethoxysilane 27326-65-4, 2-(Trimethoxysilyl)ethyl-2-pyridine 35141-30-1 84127-79-7 154260-92-1 154260-93-2 (electroless metalization of surfaces modified by, in photolithog. top-surface imaging)

L42 ANSWER 16 OF 23 HCAPLUS COPYRIGHT 2002 ACS

1993:450471 Document No. 119:50471 Characterization of films of organofunctional silanes by ToF-SIMS. Part II. Films of .gamma.-APS, AEAPS and FPS on cold-rolled steel and cold-rolled zinc substrates. van Ooij, W. J.; Sabata, A. (Armco Res. and Technol., Middletown, OH, 45044-3999, USA). Surf. Interface Anal., 20(5), 475-84 (English) 1993. CODEN: SIANDQ. ISSN: 0142-2421.

The structure, compn. and thickness of thin films of various functional silanes formed on cold-rolled steel or cold-rolled zinc surfaces were studied by high-resoln. time-of-flight SIMS and AES. The silanes were .gamma.-aminopropyltriethoxysilane, N-(2-aminoethyl)-3-aminopropyltrimethoxysilane, and perfluorinated propyltrimethoxysilane, which were deposited from dil. solns. by dipping or spin-coating. Variables were the pH of the soln. and the silane concn. The objective was to improve the understanding of the structure and orientation of silane films on metals, in view of possible applications of such coupling agents in areas where corrosion control and durable paint adhesion is required.

17 29159-37-3, .gamma.-Aminopropyltriethoxysilane homopolymer

29159-37-3, .gamma.-Aminopropyltriethoxysilane homopolymer (films, structure, compn. and thickness of, on cold-rolled steel

or cold-rolled zinc)

RN 29159-37-3 HCAPLUS 1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX CN NAME) CM 1 CRN 919-30-2 CMF C9 H23 N O3 Si OEt Eto-Si- $(CH_2)_3$ -NH<sub>2</sub> OEt 37-5 (Plastics Manufacture and Processing) CC Section cross-reference(s): 38 29159-37-3, .gamma.-Aminopropyltriethoxysilane homopolymer IT 148721-55-5 (films, structure, compn. and thickness of, on cold-rolled steel or cold-rolled zinc) 1310-73-2, Sodium hydroxide, uses IT (steel substrates treated with soln. of, characterization of silane films by time-of-flight SIMS and AES in relation to) ANSWER 17 OF 23 HCAPLUS COPYRIGHT 2002 ACS 1992:427878 Document No. 117:27878 TOF SIMS and XPS study of the interaction of hydrolyzed .gamma.-aminopropyltriethoxysilane with E-glass surfaces. Wang, D.; Jones, F. R.; Denison, P. (Sch. Mater., Univ. Sheffield, Sheffield, S10 2TZ, UK). J. Adhes. Sci. Technol., 6(1), 79-98 (English) 1992. CODEN: JATEE8. ISSN: 0169-4243. The interaction of E-glass surfaces with hydrolyzed AB .gamma.-aminopropyltriethoxysilane was studied by time-of-flight (TOF) SIMS and XPS. The SIMS spectrum consisted of 4 series of fragmentation patterns, 2 of which could be assigned to the poly(aminosilane). The remaining 2 series were consistent with the well-known fragmentation of polydimethylsiloxane. Using aq. extn. procedures, it was possible to show that the largest observable fragment increased in size after treatment with warm H2O but subsequently decreased again after hot-H2O extn. demonstrated the removal of low-mol.-wt. oligomeric species overlying a crosslinked network of graded d. The pos. Al-ion intensity remained strong, suggesting its incorporation into the silane film. XPS anal. confirmed the incorporation of aluminum ions

(formation and identification of, on glass fibers, time-of-flight SIMS and XPS study of)

29159-37-3 HCAPLUS RN

29159-37-3P

IT

from the substrate into the coating.

CN 1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 919-30-2

CMF C9 H23 N O3 Si

CC 37-6 (Plastics Manufacture and Processing)

TT 29159-37-3P

(formation and identification of, on glass fibers, time-of-flight SIMS and XPS study of)

L42 ANSWER 18 OF 23 HCAPLUS COPYRIGHT 2002 ACS

1992:175587 Document No. 116:175587 Characterization of films of organofunctional silanes by TOFSIMS and XPS. Part I. Films of N-[2-(vinylbenzylamino)ethyl]-3-aminopropyltrimethoxysilane on zinc and .gamma.-aminopropyltriethoxysilane on steel substrates. Van Ooij, W. J.; Sabata, A. (Armco Res. Technol., Middletown, OH, 45043, USA). J. Adhes. Sci. Technol., 5(10), 843-63 (English) 1991. CODEN: JATEE8. ISSN: 0169-4243.

The structure of thin films formed by N-[2-(vinylbenzylamino)-ethyl]3-aminopropyltrimethoxysilane and .gamma.-aminopropyltriethoxysilane
deposited onto mech. polished Zn or mild steel from dil. aq. solns.
were detd. using time-of-flight (TOF) SIMS and XPS. TOFSIMS gave
structural information which was highly complementary to the XPS
data. Aspects such as silane condensation and crosslinking, oxidn.
at elevated temps., the formation of metallosiloxane bonds, and
incomplete hydrolysis were detected by TOFSIMS by virtue of its high
mass resoln. and unlimited mass range. The structures of the films
were strongly dependent on the nature of the substrate,
the deposition conditions, and heat treatment of the

29159-37-3, .gamma.-Aminopropyltriethoxysilane homopolymer (films, deposited on steel, structural characterization of, by time-of-flight SIMS and XPS)

RN 29159-37-3 HCAPLUS

CN 1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 919-30-2

CMF C9 H23 N O3 Si

OEt 
$$|$$
 EtO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>  $|$  OEt

CC 38-3 (Plastics Fabrication and Uses) Section cross-reference(s): 36, 37

29159-37-3, .gamma.-Aminopropyltriethoxysilane homopolymer (films, deposited on steel, structural characterization of, by time-of-flight SIMS and XPS)

L42 ANSWER 19 OF 23 HCAPLUS COPYRIGHT 2002 ACS
1990:207946 Document No. 112:207946 Negative **pattern**-forming
composition containing polyorganosiloxane. Muramoto, Naohiro; Mine,
Katsutoshi (Toray Silicone Co., Ltd., Japan). Eur. Pat. Appl. EP
315954 A2 19890517, 17 pp. DESIGNATED STATES: R: BE, DE, FR, GB.
(English). CODEN: EPXXDW. APPLICATION: EP 1988-118581 19881108.
PRIORITY: JP 1987-282638 19871109.

A neg. pattern-forming compn. useful in producing highly ABaccurate submicron patterns having unusually high aspect ratios at superior resolns. comprises a solvent-sol. polyorganosiloxane having SiO4/2 units and at least one other organosiloxane unit contg. a high-energy-radiation-sensitive group, the polyorganosiloxane having a softening temp. greater than the The pattern-forming compn. is sensitive to room temp. electron beams and x-rays and is also sensitive to UV radiations when a sensitizer, such as 2,4-dimethylthioxanthone or 2-hydroxy-2-methylpropiophenone, is included in the compn. and can be used as a resist material in the fabrication of semiconductor devices. The pattern formed from the compn. is ideally used as a mask highly resistant to the dry etching of an underlying org. material, such as an org. insulating material or an org. planarizing layer as used in a multilayer resist

1T 4420-74-0, 3-Mercaptopropyltrimethoxysilane (reaction of, in prepg. polyorganosiloxanes as radiation resists)

RN 4420-74-0 HCAPLUS

CN 1-Propanethiol, 3-(trimethoxysilyl)- (7CI, 8CI, 9CI) (CA INDEX NAME)

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IC
     ICM G03F007-10
CC
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and
     Other Reprographic Processes)
     Section cross-reference(s): 76
     polyorganosiloxane radiation resist semiconductor device;
ST
     x ray resist polyorganosiloxane; electron beam
     resist polyorganosiloxane; photoresist UV
     polyorganosiloxane semiconductor device
IT
     Siloxanes and Silicones, uses and miscellaneous
       (as radiation resists for forming dry-etching-resistant
        patterns for semiconductor device fabrication)
     Semiconductor devices
IT
        (polyorganosiloxane radiation resists for fabrication
        of)
     Resists
IT
        (electron-beam, polyorganosiloxanes as)
     Resists
IT
        (photo-, UV, polyorganosiloxanes as)
     Resists
IT
        (x-ray, polyorganosiloxanes as)
IT
     75-77-4, Trimethylchlorosilane, reactions
                                                 78-10-4,
                         682-01-9, Tetrapropyloxysilane
                                                          999-97-3,
     Tetraethoxysilane
     Hexamethyldisilazane
                            1719-57-9, 1-Chloromethyldimethylchlorosilane
     1719-58-0, Dimethylvinylchlorosilane
                                            1825-61-2
                                                        2530-85-0,
     3-Methacryloxypropyltrimethoxysilane
                                            2768-02-7,
                             3634-56-8, Dimethylisopropylchlorosilane
     Vinyltrimethoxysilane
     4420-74-0, 3-Mercaptopropyltrimethoxysilane
                                                   7446-81-3,
                       7691-02-3, 1,3-Divinyl-1,1,3,3-
     Sodium acrylate
     tetramethyldisilazane
                             14346-37-3, Isopropyltrimethoxysilane
     15933-59-2, 1,3-Dihydrogen-1,1,3,3-tetramethyldisilazane
                 16753-62-1, Methylvinyldimethoxysilane
     16546-47-7
                                                           18143-33-4
     18162-48-6, tert-Butyldimethylchlorosilane
                                                  103460-81-7,
     p-(1-Chloromethyl) phenethyltrichlorosilane
                                                  116778-00-8,
     1,3-Bis(3-glycidoxypropyl)-1,1,3,3-tetramethyldisilazane
        (reaction of, in prepq. polyorganosiloxanes as radiation
        resists)
    ANSWER 20 OF 23 HCAPLUS COPYRIGHT 2002 ACS
1989:163595 Document No. 110:163595 Photosensitive polyamic acid
     compositions. (Shin-Etsu Chemical Industry Co., Ltd., Japan).
     Kokai Tokkyo Koho JP 60088939 A2 19850518 Showa, 20 pp. (Japanese).
```

CODEN: JKXXAF. APPLICATION: JP 1983-197129 19831021.

GI

The claimed photosensitive compns. contain (1) a polyamic acid AB obtained by reaction of (CH2:CHZd)a(H2NZ1)bR1cSiO(4-a-b-c)/c (0< a; b .ltoreq. 3; 0 .ltoreq. c .ltoreq. 2; a + b + c .ltoreq. 4; d = 0, 1; Z, Z1 = divalent org. moiety), I (<math>Z2 = tetravalent org. moiety), and H2NZ3NH2 or H2NZ4Z5(NH2)2 (Z3 = divalent org. moiety; Z4 = SO2, CO' Z5 = trivalent org. moiety), (2) a mercapto compd. contg. .gtoreq.2 SH groups: less than equiv. with respect to the CH2CHZd groups of the above Si compds.), and (3) a photosensitizer [0-5 wt. parts/100 parts solids on (1) + (2)]. The compns. exhibit excellent storage stability, good resolving power, and good adhesion with substrates. The photosensitive compns. are esp. useful in forming heat-resistant polyimide insulator patterns.

108315-45-3 IT

CN

(photosensitive polyamic acid compns. contg.)

108315-45-3 HCAPLUS RN

1-Propanethiol, 3,3',3'',3'''-(1,3,5,7-tetramethyl-1,3,5,7tetrasiloxanetetrayl)tetrakis- (9CI) (CA INDEX NAME)

IC ICM G03C001-71

ICS C08G075-12; G03F007-10

CC 74-4 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

IT Semiconductor devices

(fabrication of, resist compns. for)

IT Polyimides, uses and miscellaneous

(insulator patterns of, prepn. of)

IT Polyamic acids

(photosensitive compns. contg., polyimide insulator patterns from)

IT Resists

(photo-, siloxane deriv. copolymers as)
IT 80-43-3 18001-52-0 22499-12-3 108315-45-3
(photosensitive polyamic acid compns. contg.)

L42 ANSWER 21 OF 23 HCAPLUS COPYRIGHT 2002 ACS

1988:561967 Document No. 109:161967 Selective metallization process and additive method for manufacturing printed circuit boards. Ferrier, Donald R.; Larson, Gary B. (MacDermid, Inc., USA). U.S. US 4748104 A 19880531, 7 pp. (English). CODEN: USXXAM. APPLICATION: US 1986-929595 19861110.

AB Processes for selective metalization of a substrate are described in which a substrate **patterned** in a desired manner with a **resist** is treated with a conditioning agent (which increases the adsorption on exposed substrate areas of subsequently applied

catalytic species), an adjuvant (which decreases the receptivity of the resist for the catalytic species), and a deactivating agent (under conditions which also decrease the receptivity of the resist for the catalytic species) preparatory to catalytic activation and electroless deposit of metal (e.g., Cu) flash. By proceeding in this manner, adherent flash metal deposit is achieved on non-resist areas without encountering significant plating on resist areas. Following metal flash deposit, the resist can be easily and cleanly stripped and addnl. metal then built up on the flash metal. The adjuvant may be an amine; the conditioning agent may be an organosilane; and the deactivating agent may comprise an aq. alk. soln. 1760-24-3

(in selective metalization for printed circuit board manuf.) 1760-24-3 HCAPLUS

CN 1,2-Ethanediamine, N-[3-(trimethoxysilyl)propyl]- (9CI) (CA INDEX NAME)

OMe 
$$\mid$$
 MeO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH-CH<sub>2</sub>-CH<sub>2</sub>-NH<sub>2</sub>  $\mid$  OMe

IC ICM G03C005-00

NCL 430315000

IT

RN

CC 76-14 (Electric Phenomena)

7440-50-8, Copper, uses and miscellaneous
 (electroless deposition of, pretreatment of
 substrates for selective metalization by, in printed
 circuit board manuf.)

TT 78-90-0, 1,2-Propanediamine 97-84-7, N,N,N',N'-Tetramethyl-1,3-butanediamine 107-15-3, Ethylenediamine, uses and miscellaneous 111-40-0, Diethylene triamine 497-19-8, Sodium carbonate, uses and miscellaneous 919-30-2, .gamma.-Aminopropyltriethoxysilane 1760-24-3 3010-24-0 116711-11-6

(in selective metalization for printed circuit board manuf.)

L42 ANSWER 22 OF 23 HCAPLUS COPYRIGHT 2002 ACS

1987:76145 Document No. 106:76145 Lift-off processes by multilayer resists. Ono, Yoichi (Seiko Epson K. K., Japan). Jpn.
Kokai Tokkyo Koho JP 61170738 A2 19860801 Showa, 4 pp. (Japanese).
CODEN: JKXXAF. APPLICATION: JP 1985-11859 19850125.

AB In lift-off processes with multilayer resists, the surfaces of underlayer resists are pretreated with silane coupling agents to prevent dissoln. of the resists in each other. Thus, a pos.-type photoresist (thickness .apprx.2 .mu.) was applied to a Si wafer and prebaked at 80.degree. for 20 min, then exposed all over the surface so as to make the exposure 2 times of the upper

photoresist, surface treated with a 0.1

vol.% aq. SH-6020 (silane coupling agent), dried at 80.degree. for 10 min, then the same <code>resist</code> (thickness 0.8 .mu.) was applied again, prebaked at 100.degree. for 20 min, exposed to light through a photomask, developed to form an over-hung shape, then a 2000-.ANG. Al film was formed on it, and finally the <code>resist</code> was peeled off to obtain an Al <code>pattern</code>.

IT 1760-24-3, SH-6020

(coupling agent, in lift-off process with multilayer
resists)

RN 1760-24-3 HCAPLUS

CN 1,2-Ethanediamine, N-[3-(trimethoxysilyl)propyl]- (9CI) (CA INDEX NAME)

OMe 
$$\mid$$
 MeO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH-CH<sub>2</sub>-CH<sub>2</sub>-NH<sub>2</sub>  $\mid$  OMe

IC ICM G03C005-00 ICS G03C001-00; G03F007-00

ICA H01L021-30

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST multilayer resist lift off process; silane coupling agent multilayer resist

IT Resists

(lift-off process with multilayer, silane coupling agent in)
1760-24-3, SH-6020 61417-49-0, TTS 106716-89-6
(coupling agent, in lift-off process with multilayer resists)

L42 ANSWER 23 OF 23 HCAPLUS COPYRIGHT 2002 ACS

1967:86774 Document No. 66:86774 Corrosion-resistant coatings for metal. Morehouse, Edward L.; Pines, Arthur N. (Union Carbide Corp.). Ger. DE 1233522 19670202, 10 pp. (German). CODEN: GWXXAW. PRIORITY: US 19570719.

The prepn. of an anticorrosive coating for metal is described. Thus, 50 g. H2O was added to a mixt. of 200 g. .gamma.—
aminopropyltriethoxysilane (I) and 130 g. abs. EtOH (II) over 15-20 min. and the mixt. was heated under reflux for 30 min. This soln. was dild. with II to give 1 part resin/5 parts II. A Cu strip was partially coated with the dild. soln. and air-dried. The strip was heated for 4 hrs. at 250.degree. in an air oven. The treated surface was bright, while the untreated portion was oxidized and dark. A 5% alc. soln. of I homopolymer (III) or of III-PhSi(OEt)3 copolymer or of [H2N(CH2)3SiO1.5]0.5(PhSiO1.5)0.4[Me2SiO]0.1 (IV) can be used to coat Cu, Al, or Ag in the case of IV.

```
29159-37-3
TT
        (coatings of, on metal, corrosion-resistant)
     29159-37-3 HCAPLUS
RN
     1-Propanamine, 3-(triethoxysilyl)-, homopolymer (9CI) (CA INDEX
CN
     NAME)
     CM
     CRN
          919-30-2
     CMF
          C9 H23 N O3 Si
     OEt.
Eto-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>
     OEt
IC
     C09D
CC
     42 (Coatings, Inks, and Related Products)
IT
                3037-72-7
                             3069-30-5
                                         13497-18-2
     919-30-2
     16099-52-8 29159-37-3
                              31392-08-2
                                           31392-09-3
     31392-10-6
                  31392-11-7
                                31392-12-8
                                             31392-15-1
                                                           31392-25-3
     31474-05-2
        (coatings of, on metal, corrosion-resistant)
=> d 143 1-8 cbib abs hitstr hitind
     ANSWER 1 OF 8 HCAPLUS COPYRIGHT 2002 ACS
2001:762370 Document No. 135:325313 Manufacture of optical disk master
     with high resolution pattern. Sakamizu, Toshio;
     Shiraishi, Hiroshi (Hitachi Ltd., Japan). Jpn. Kokai Tokkyo Koho JP
     2001291288 A2 20011019, 6 pp. (Japanese). CODEN: JKXXAF.
     APPLICATION: JP 2000-107998 20000405.
     The title optical disk master manuf. comprises (1) a process to
AB
     treat a substrate surface with a
     nonbase-generating type org. silicon compd. contg. a hydrolysis
     group(s) or a vinyl polymer contg. a hydrolysis group(s), (2) a
     process to form a chem. amplification resist compn. layer
     on the substrate, (3) a process to irradiate the
     photoresist layer to form a latent pit pattern,
     and (4) a process to develop the pit pattern. The manuf.
     produces the optical disk master with high throughput.
     919-30-2, 3-Aminopropyltriethoxysilane 2530-83-8,
IT
     3-Glycidoxypropyltrimethoxysilane
        (for surface treatment of optical
        disk substrate)
RN
     919-30-2 HCAPLUS
    1-Propanamine, 3-(triethoxysilyl)- (9CI) (CA INDEX NAME)
CN
```

OEt 
$$|$$
 EtO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>  $|$  OEt

RN 2530-83-8 HCAPLUS

CN Silane, trimethoxy[3-(oxiranylmethoxy)propyl]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} O & OMe \\ \hline \\ CH_2-O-(CH_2)_3-Si-OMe \\ \hline \\ OMe \\ \end{array}$$

IC ICM G11B007-26

CC 74-12 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST optical disk master manuf surface treatment

silane coupling agent

IT Photoresists

(chem. amplification; manuf. of optical disk master with high resoln. pattern)

IT Optical disks

Photolithography

(manuf. of optical disk master with high resoln. pattern

75-94-5, Vinyltrichlorosilane 919-30-2,
3-Aminopropyltriethoxysilane 2530-83-8,
3-Glycidoxypropyltrimethoxysilane 2530-85-0, 3Methacryloxypropyltrimethoxysilane
(for surface treatment of optical

disk substrate)

IT 9003-17-2, Polybutadiene

(partially epoxylated; for surface treatment of optical disk substrate)

L43 ANSWER 2 OF 8 HCAPLUS COPYRIGHT 2002 ACS

2001:213015 Document No. 135:38836 Large-area MEMS fabrication with thick SU-8 photoresist applied to an x-ray image sensor array. Daniel, Jurgen H.; Krusor, Brent S.; Apte, Raj B.; Street, Robert A.; Goredema, Adela; McCallum, Jason; Boils-Boissier, Daniele C.; Kazmaier, Peter M. (Xerox Palo Alto Research Center, Palo Alto, CA, 94304, USA). Proceedings of SPIE-The International Society for Optical Engineering, 4174 (Micromachining and Microfabrication Process Technology VI), 40-48 (English) 2000. CODEN: PSISDG. ISSN:

0277-786X. Publisher: SPIE-The International Society for Optical Engineering.

AB MEMS fabrication on large area substrates is promising for novel system concepts, but processes based on cryst. silicon cannot be used. Polymeric materials such as the thick photoresist SU-8 are more appropriate for this purpose because their processing can be scaled to large areas. An x-ray image sensor array based on amorphous silicon on glass substrates was taken as an examples to apply the SU-8 microfabrication technique. The authors anticipate an improved performance of the imager. The resoln. of the x-ray imager is expected to increase by patterning the x-ray conversion screen into cells, which match the sensor pixels of the imager, and SU-8 defines these cells. Furthermore, SU-8 functions as a thick interlayer dielec. with decrease the electronic noise and result in a higher sensitivity of the imager. The fabrication process with SU-8 will be described and challenging issues, esp. when considering large area substrates, will be discussed.

IT 2530-83-8, 3-Glycidoxypropyltrimethoxysilane

(adhesion promoter; microfabrication of x-ray image sensor array using thick **photoresist** SU-8 process and application of SR-U as interlayer dielec.)

RN 2530-83-8 HCAPLUS

CN Silane, trimethoxy[3-(oxiranylmethoxy)propyl]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} \text{OMe} & \text{OMe} \\ \mid & \mid \\ \text{CH}_2\text{-O-(CH}_2)_3\text{-Si-OMe} \\ \mid & \mid \\ \text{OMe} \end{array}$$

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST microfabrication x ray image sensor array thick SU8 photoresist

IT Radiography

(digital; microfabrication of amorphous Si on glass substrates x-ray image sensor array using thick **photoresist** SU-8 process)

IT Micromachining

Photoresists

(microfabrication of amorphous Si on glass substrates x-ray image sensor array using thick **photoresist** SU-8 process)

IT Electric insulators

Electrodeposition

(microfabrication of x-ray image sensor array using thick **photoresist** SU-8 process and application of SR-U as interlayer dielec.)

- IT Optical imaging sensors
   (x-ray; microfabrication of amorphous Si on glass substrates
   x-ray image sensor array using thick photoresist SU-8
   process)
- 2530-83-8, 3-Glycidoxypropyltrimethoxysilane (adhesion promoter; microfabrication of x-ray image sensor array using thick **photoresist** SU-8 process and application of SR-U as interlayer dielec.)

- IT 2551-62-4, Sulfur hexafluoride (plasma; microfabrication of x-ray image sensor array using thick **photoresist** SU-8 process and application of SR-U as interlayer dielec.)
- 2000:682637 Document No. 134:11384 Selective Deposition in Layer-by-Layer Assembly: Functional Graft Copolymers as Molecular Templates. Jiang, Xueping; Hammond, Paula T. (Department of Chemical Engineering, Massachusetts Institute of Technology, Cambridge, MA, 02139, USA). Langmuir, 16(22), 8501-8509 (English) 2000. CODEN: LANGD5. ISSN: 0743-7463. Publisher: American

ANSWER 3 OF 8 HCAPLUS COPYRIGHT 2002 ACS

L43

Chemical Society.

AB A new approach to creating **patterned** composite structures with layer-by-layer assembled thin films is introduced. In this approach, the top surfaces of layer-by-layer films are chem.

patterned by the direct stamping of functional polymers on the surface. The resulting pattern is then used as a template for the further deposition of materials on the surface. This paper illustrates this concept through the patterning of an oligoethylene glycol resist atop a polyelectrolyte multilayer surface. The resist layer is created by stamping an oligoethylene oxide-maleic anhydride graft copolymer onto the top polyamine surface. Further deposition of polyions results in the formation of a patterned polymer film atop the original multilayer thin film. When a patterned polyelectrolyte film is used as the base layer or substrate in this process, a second pattern can be stamped atop the original patterned array. Subsequent selective adsorption of polymers yields a second level of microstructures, illustrating the concept of multiple level patterning for layer-by-layer thin films; this concept can be broadened to include selective deposition of numerous materials on top of multilayer polymer systems. 13822-56-5, Aminopropyltrimethoxysilane

IT (patterning oligoethylene oxide-maleic anhydride graft copolymer on aminosilane SAM model surface)

RN 13822-56-5 HCAPLUS

1-Propanamine, 3-(trimethoxysilyl)- (9CI) (CA INDEX NAME)

$$\begin{array}{c} \text{OMe} \\ | \\ \text{MeO-Si-} (\text{CH}_2)_3 - \text{NH}_2 \\ | \\ \text{OMe} \end{array}$$

CN

74-5 (Radiation Chemistry, Photochemistry, and Photographic and CC Other Reprographic Processes)

polyelectrolyte multilayer patterning graft copolymer mol STtemplate; oliqoethylene oxide maleic anhydride graft copolymer template adsorption polyelectrolyte; lithog patterning graft copolymer mol template adsorption polyelectrolyte multilayer

IT Chemisorption IR spectra Multilayers

Physisorption

Polyelectrolytes

Resists

(lithog. by printing surface of multilayer film with oligoethylene oxide-maleic anhydride graft copolymer resist to template subsequent selective multilayer adsorption of polyelectrolytes)

Self-assembled monolayers IT

(patterning complex microstructures by stamping oligoethylene oxide-maleic anhydride graft copolymer on patterned surfaces)

IT Lithography (use of functional template from oligoethylene oxide-maleic anhydride graft copolymer for chem. patterning of layer-by-layer film surfaces) 112311-92-9, Maleic anhydride-polyethylene glycol allyl IT 215512-58-6 methyl ether copolymer (lithog. by printing surface of multilayer film with oligoethylene oxide-maleic anhydride graft copolymer resist to template subsequent selective multilayer adsorption of polyelectrolytes) IT 130727-41-2 (patterning complex microstructures by stamping oliqoethylene oxide-maleic anhydride graft copolymer on patterned surfaces) 13822-56-5, Aminopropyltrimethoxysilane IT (patterning oligoethylene oxide-maleic anhydride graft copolymer on aminosilane SAM model surface) 9003-04-7, Poly(acrylic acid) sodium salt IT 9080-79-9, Sodium poly(styrenesulfonate) 26062-79-3, Polydiallyldimethylammonium chloride 26913-06-4, Poly[imino(1,2-ethanediyl)] (polyelectrolyte; lithog. by printing surface of multilayer film with oliquethylene oxide-maleic anhydride graft copolymer resist to template subsequent selective multilayer adsorption of polyelectrolytes) 9016-00-6, Poly(dimethylsiloxane) 31900-57-9, IT Poly(dimethylsiloxane) (stamp; lithog. by printing surface of multilayer film with oligoethylene oxide-maleic anhydride graft copolymer resist to template subsequent selective multilayer adsorption of polyelectrolytes) 7440-21-3, Silicon, processes 7440-57-5, Gold, IT processes 9003-53-6, Polystyrene (substrate; lithog. by printing surface of multilayer film with oligoethylene oxide-maleic anhydride graft copolymer resist to template subsequent selective multilayer adsorption of polyelectrolytes) ANSWER 4 OF 8 HCAPLUS COPYRIGHT 2002 ACS Document No. 130:318478 Using Elastomeric Membranes as Dry 1999:183139 Resists and for Dry Lift-Off. Jackman, Rebecca J.; Duffy, David C.; Cherniavskaya, Oksana; Whitesides, George M. (Department of Chemistry and Chemical Biology, Harvard University, Cambridge, Langmuir, 15(8), 2973-2984 (English) 1999. MA, 02138, USA). ISSN: 0743-7463. Publisher: American Chemical Society. Elastomeric membranes that contained regular arrays of well-defined AB holes were formed by spin-coating a prepolymer onto a photolithog. These membranes were used as dry resists defined master. or as masks in dry lift-off to produce simple features .gtoreq.5 .mu.m on both planar and nonplanar surfaces. These procedures were dry because the membranes conformed and sealed reversibly to

surfaces: no solvent was required either to deposit the membrane or

to remove it from the substrate. A variety of materials, some of which would be difficult to pattern using conventional methods, were patterned using this technique. These materials included metals, sol-gels, hydrogels, biol. macromols., and organometallic mols. The membranes were used in sequential, dry-lift off steps to produce structures with greater complexity than those generated with a single membrane.

2530-83-8, [3-(Glycidyloxy)propyl]trimethoxysilane IT

(micropatterning of org. modified silicate formed by sol-gel precursor using polydimethylsioloxane membrane as dry lithog. resists)

RN 2530-83-8 HCAPLUS

Silane, trimethoxy[3-(oxiranylmethoxy)propyl]- (9CI) (CA INDEX CN NAME)

$$\begin{array}{c|c} \text{OMe} & \text{OMe} \\ | & \\ \text{CH}_2\text{--O-(CH}_2)_3\text{--Si--OMe} \\ | & \\ \text{OMe} \end{array}$$

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

elastomeric membrane dry resist lift off lithog ST

Membranes, nonbiological IT

(elastomeric; micropatterning using elastomeric membranes as dry lithog. resists and for dry lift-off)

Ceramers IT

Sol-gel processing

(micropatterning of org. modified silicate formed by sol-gel precursor using polydimethylsioloxane membrane as dry lithog. resists)

Resists IT

> (micropatterning using elastomeric membranes as dry lithog. resists and for dry lift-off)

IT Gelatins, processes

> (micropatterns of gelatin using polydimethylsioloxane membrane as dry lithoq. resists)

IT Electron beams

> (micropatterns of gold fabricated by electron-beam evapn. using polydimethylsioloxane membrane as dry lithog. resists)

IT Electrodeposition

(micropatterns of silver deposited by electroplating using polydimethylsioloxane membrane as dry lithog. resists) 144-62-7, Oxalic acid, processes 2530-83-8,

IT

[3-(Glycidyloxy)propyl]trimethoxysilane (micropatterning of org. modified silicate formed by sol-gel precursor using polydimethylsioloxane membrane as dry lithog. resists)

- IT 7440-22-4, Silver, processes

  (micropatterning of silver by electroplating using polydimethylsioloxane membrane as dry lithog. resists)
- 9016-00-6, Polydimethylsiloxane 31900-57-9, Polydimethylsiloxane (micropatterning using elastomeric membranes as dry lithog. resists and for dry lift-off)
- L43 ANSWER 5 OF 8 HCAPLUS COPYRIGHT 2002 ACS
- 1998:439857 Document No. 129:181988 One mask process for stress-buffer and passivation applications using photosensitive benzocyclobutene. Strandjord, A. J. G.; Rogers, W. B.; Ida, Y.; Shiau, S.; Moyer, E. S.; Scheck, D. M.; Devellis, R. R.; Garrou, P. E. (Microelectronics Research and Development, The Dow Chemical Company, Midland, MI, 48674, USA). IEMT/IMC Symposium, 1st, Sonic City-Omiya, Japan, Apr. 16-18, 1997, 261-266. SHM: The Microelectronics Society: Tokyo, Japan. (English) 1997. CODEN: 66JZAY.
- Large die such as SRAMs and DRAMs are subject to significant AB stresses resulting from the differences in the coeff. of thermal expansion (CTE mismatch) of the die and packaging materials. use of polymers as a buffering medium between these two materials has been widely practiced in the industry to increase the reliability of these devices. This layer must be patterned to allow elec. connection to the bond pads (50 - 250 .mu.m openings). In many of the newer devices, smaller windows (<10 .mu.m) must also be opened up to allow access to other structures, such as "fuse links", which are used to reroute or reconfigure inactive circuits. This need for higher resoln. polymer layers, as well as the desire for shorter processing cycles, has moved the industry away from the use of wet etch polymers toward Traditional wafer level photosensitive materials. processing of memory die has required a two mask operation for patterning features in the underlying primary passivation layer (commonly Si3N4) followed by coating and patterning of the polymer stress-buffer layer. A one mask manufg. process has been developed using photosensitive benzocyclobutene (Photo-BCB) for opening up features (bond pads and fuse links) in both layers using the polymer layer as a permanent etch mask for the Si3N4. This results in a significant redn. in both the no. of processing steps and the processing time, vs. the traditional two mask process, and therefore, significantly lowers the cost of ownership for passivation/stress-buffer layers. paper, we will discuss the Photo-BCB processing steps and compare them to the steps in a two mask passivation/stress-buffer processes. IT 919-30-2, 3-Aminopropyltriethoxysilane

(adhesion promoter; photolithog. one mask process for stress-buffer and passivation applications using photosensitive

benzocyclobutene) RN 919-30-2 HCAPLUS

CN 1-Propanamine, 3-(triethoxysilyl)- (9CI) (CA INDEX NAME)

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 76

ST memory die wafer level processing; photolithog single mask process Cyclotene

IT Photoresists

IT

(Cyclotene; photolithog. one mask process for stress-buffer and passivation applications using photosensitive benzocyclobutene) 919-30-2, 3-Aminopropyltriethoxysilane

(adhesion promoter; photolithog. one mask process for stress-buffer and passivation applications using photosensitive benzocyclobutene)

L43 ANSWER 6 OF 8 HCAPLUS COPYRIGHT 2002 ACS
1992:13389 Document No. 116:13389 Surface treatment
 of substrates with resist patterns.
 Inamoto, Tadaki (Canon K. K., Japan). Jpn. Kokai Tokkyo Koho JP
 03184051 A2 19910812 Heisei, 8 pp. (Japanese). CODEN: JKXXAF.
 APPLICATION: JP 1989-322750 19891214.

AB The treatment consists of treating substrate

The treatment consists of treating substrate having resist pattern with 0.01-10 wt.% solns. of silane coupling agent in solvents that do not dissolve or deform the pattern, and subsequent removal of the coupling agent with the same solvent. Typically the resist pattern is made from pos.-working photoresist contq. quinonediazides and alkali-sol. resins. This treatment is useful in working in micro scale using the pattern as mold, e.g. in manuf. of ink head for ink-jet printing, by ensuring close contact of molding agent with the resist pattern Thus, a Si wafer on which 2 heat-generating elements for ink-jet head were mounted was appropriately patterned by photolithog. using Ozatec R225 ( photoresist). The pattern surface was sprayed for 1 min with 0.01% soln. of A187 (silane coupling agent), and for 3 min with water. The **pattern** was potted with a photocurable bisphenol A epoxy resin, treated with

1,1,1-trichloroethane, and sonicated in 3% NaOH for 5 min to remove the pos. photoresist pattern, to obtain an

ink-jet head with 2 orifices and 2 heat-generating elements. No defect was obsd. in the head after using for printing 50,000 copies

$$CH_2-O-(CH_2)_3-Si-OMe$$
OMe
OMe
OMe

IC ICM G03F007-40 ICS B41J002-16

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST resist pattern surface treatment molding; silane coupling

treatment molding; silane coupling agent resist
treatment; ink jet printing head manuf

IT Coupling agents

(silane, treatment with, of resist pattern used for micromolding)

IT Printing, nonimpact

(ink-jet, manuf. of head for, micromolding using resist pattern in, surface treatment of resist pattern in)

IT Resists

(photo-, pattern of, surface treatment of, for micromolding, silane coupling agents for)

IT 110158-67-3, Ozatec R225

(pattern of, surface treatment of,

in micromolding process, silane coupling agents for)

IT 2530-83-8, A187

(treatment of resist pattern with, in micromolding process)

L43 ANSWER 7 OF 8 HCAPLUS COPYRIGHT 2002 ACS

1987:468170 Document No. 107:68170 Preparation of negative- and positive-working **photoresists** resistive to dry etching.

Iwamoto, Masao (Toray Industries, Inc., Japan). Jpn. Kokai Tokkyo Koho JP 61270755 A2 19861201 Showa, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1985-112084 19850527.

AB The title **photoresists** are comprised of a photosensitive layer contg. a quinoneazide deriv. and a silicone rubber layer. A typical processing method includes total exposure of the **photoresists** to decomp. 5-60 mol% of the quinoneazide

deriv., imagewise exposure to decomp. .gtoreq.70 mol% of the quinoneazide deriv. (the order of the exposures may be reversed), treatment with a base, and development. Alternately, the photoresists may be imagewise exposed to decomp. 5-60% of quinoneazide deriv., treatment with a base, and development. The method provides resist patterns highly resistive to dry etching, with wide selection of materials. Esp. high photosensitivity is obtained for formation of pos. patterns. The method is useful in manuf. of integrated elec. circuit patterns and also for O-plasma etching of surface of plastics. Thus, a Si wafer was coated with a 2-.mu. layer contq. a partial (47%) ester of a phenol-HCHO novolak resin with naphthoquinone-1-2-diazido-5-sulfonic acid 100, 4,4'-diphenylmethanediisocyanate 30, and dibutyltin dilaurate 0.2 parts, baked at 120.degree. for 2 min, and coated with 1-.mu. layer contg. dimethylpolysiloxane (mol. wt. 25,000; terminal OH) 100, vinyltris(methylethylketoxime)silane 8, dibutyltin diacetate 0.2, and .gamma.-aminopropyltrimethoxysilane 1 part. Total exposure with a metal halide lamp (11 mW/cm2) for 6 s to decomp. 18% of the quinoneazide deriv., exposure through a neg. pattern (60 s), treatment in a 90:5:5:0.1 Isopar H-butylcarbitol-Et Cellosolve-ethanolamine, development in a 2:4:4 Isopar H-butylcarbitol-ethylcarbitol, and rubbing with soft pad to remove the exposed part of the silicone rubber gave a pattern, which was highly resistant to 0 plasma dry etching. 13822-56-5, .gamma.-Aminopropyltrimethoxysilane (in prepn. of silicone rubber top layer for photoresist contq. quinonediazide deriv. layer) 13822-56-5 HCAPLUS 1-Propanamine, 3-(trimethoxysilyl)- (9CI) (CA INDEX NAME) OMe OMe

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MeO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>
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IT

RN

CN

IC ICM G03C005-24

G03C001-00; G03F007-00; H01L021-30 ICS

74-5 (Radiation Chemistry, Photochemistry, and Photographic and CC Other Reprographic Processes)

photoresist silicone rubber etching resistant;
quinonediazide silicone photoresist etching resistant ST

Rubber, silicone, uses and miscellaneous IT (photoresists contg. layer of, for dry

etching-resistant images, preexposure in processing of)

Alkanes, uses and miscellaneous IT

(C9-12-iso-, photoresist treatment in soln. contq., for photosensitive layer contg. quinonediazide deriv. and silicone rubber)

(photo-, contg. quinonediazide deriv. layer and silicone rubber layer for dry etching-resistant images, preexposure in processing

of)

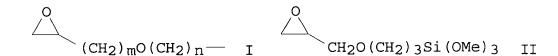
IT 2224-33-1, Vinyltris(methylethylketoxime)silane 13822-56-5, .gamma.-Aminopropyltrimethoxysilane (in prepn. of silicone rubber top layer for photoresist contg. quinonediazide deriv. layer)

TT 77-58-7, Dibutyltin dilaurate 101-68-8 27156-72-5 (photoresist contg. silicone rubber layer and layer contg., for dry etching-resistant images)

IT 110-80-5 111-90-0 112-34-5 141-43-5, uses and miscellaneous (photoresist treatment in soln. contg., for photosensitive layer contg. quinonediazide deriv. and silicone rubber)

L43 ANSWER 8 OF 8 HCAPLUS COPYRIGHT 2002 ACS
1986:59419 Document No. 104:59419 Resist formation. Fujii,
Tsuneo (Daikin Kogyo Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP
60138546 A2 19850723 Showa, 5 pp. (Japanese). CODEN: JKXXAF.
APPLICATION: JP 1983-248533 19831227.

GΙ



A resist formation method consists of treating a AB substrate surface with a silane coupling agent having amino, thiol, vinyl, or ethylene oxide groups and then forming a resist layer consisting of a fluoroalkyl acrylate polymer. Preferable silane coupling agents have .gtoreq.1 of the H atoms attached to the Si atom (of a silane, silathiane or siloxane) substituted by H2N(CH2)n, H2N(CH2)nNH(CH2)n, H2NCONH(CH2)n, HS(CH2)n, H2C:CH, or I group (n, m = 1-5) and .gtoreq.1 of the other H atoms by C1-5 alkoxy, C2-6 methoxyalkoxy, or Cl and (optionally) .gtoreq.1 of the other H atoms by C1-5 alkyl. Useful compds. are HiNC3H6Si(OEt)3, H2NC2H4NHC3H6Si(OMe)3, H2NCONHC3H6Si(OEt)3, HSC3H6Si(OMe)3, H2C:CHSi(OMe)3, H2C:CHSi(OEt)3, H2C:CHSi(OC2H4OMe)3, H2C:CHSiCl3, and II. Typically the fluoroacrylate polymer is selected from homopolymers and copolymers of monomers H2C:CRCO2ZR1 (R = Me, Et, halomethyl, haloethyl, halo, H; R1 = C1-15 fluoroalkyl; Z = C1-6 divalent hydrocarbon group). The resists have high adhesivion to substrate surfaces. Thus, a soln. contq. .gamma.-aminopropyltriethoxysilane 5, H2O 5,

and EtOH 90 parts was spin-coated on a Si wafer and prebaked. mixt. of 2,2,3,4,4,4-hexafluorobutyl methacrylate 99 and glycidyl methacrylate 1 part was polymd. to obtain a copolymer having a wt. av. mol. wt. of 1.2 .times. 106. A MEK soln. of the copolymer was coated on the precoated wafer, prebaked to form a 0.4-.mu. layer, imagewise irradiated by an electron beam, and developed with a 1.5:98.5 Me iso-Bu ketone-iso-PrOH mixt. No defects due to sepn. of the resist layer was obsd. in .gtoreq.2-.mu. line and space patterns 919-30-2 (coupling agent, silicon wafer surfaces pretreated with, for coating of fluoroalkyl acrylate copolymer electron-beam resist layers for fabrication of semiconductor devices)

RN 919-30-2 HCAPLUS

IT

1-Propanamine, 3-(triethoxysilyl)- (9CI) (CA INDEX NAME) CN

$$\begin{array}{c} \text{OEt} \\ \mid \\ \text{EtO-Si-} (\text{CH}_2)_3 - \text{NH}_2 \\ \mid \\ \text{OEt} \end{array}$$

IC ICM G03C001-74

B05D005-12; B05D007-24; G03F007-16; H01L021-30

74-5 (Radiation Chemistry, Photochemistry, and Photographic and CC Other Reprographic Processes)

photoresist fluoroalkyl acrylate polymer adhesion; silane ST coupling agent fluoropolymr resist

Semiconductor devices IT

> (coating of fluoroalkyl acrylate copolymer electron-beam resist layer on silicon wafer surfaces pretreated with silane coupling agent for fabrication of)

IT

(electron-beam, fluoroalkyl acrylate copolymer, coating of, on silicon wafer surfaces pretreated

with silane coupling agent)

919-30-2 IT

(coupling agent, silicon wafer surfaces pretreated with, for coating of fluoroalkyl acrylate copolymer electron-beam resist layers for fabrication of semiconductor devices)

67-63-0, uses and miscellaneous TT 108-10-1 (developer contg., for fluoroalkyl acrylate copolymer electron-beam resist coated on silicon wafers pretreated with silane coupling agent)

142-82-5, uses and miscellaneous IT 108-83-8 (developer, for fluoroalkyl acrylate copolymer electron-beam resist coated on silicon wafers pretreated with silane coupling agent)

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66396-71-2
IT
                  86904-68-9
                               87458-36-4
                                             99936-06-8
        (electron-beam resist, coating of, on silicon
        wafers pretreated with silane coupling agent
        for fabrication of semiconductor devices)
IT
     7440-21-3, uses and miscellaneous
        (wafers, treated with silane coupling agent
        for coating of fluoroalkyl acrylate copolymer electron-beam
        resist layers for fabrication of semiconductor devices)
=> d his 144-
     (FILE 'HCAPLUS' ENTERED AT 17:19:25 ON 15 MAY 2002)
     FILE 'REGISTRY' ENTERED AT 18:05:09 ON 15 MAY 2002
     FILE 'HCAPLUS' ENTERED AT 18:06:27 ON 15 MAY 2002
          15652 S PHOTOLITHO? OR PHOTO(2A)LITHO?
L44
L45
             70 S L34 AND L44
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9 S L45 AND L7 AND L19 L48 13 S L45 AND L7 AND L15 L49

7 S L45 AND L15 AND L19

L50 11 S (L47 OR L48 OR L49) NOT (L42 OR L43)

55 S L45 AND (L7 OR L15 OR L19)

=> d 150 1-11 cbib abs hitstr hitind

L46

L47

L50 ANSWER 1 OF 11 HCAPLUS COPYRIGHT 2002 ACS 2001:721614 Document No. 135:249448 Photomasking of photosensitive polyimides with organometallic compounds for photolithography on silicon chips. Munoz Pascual, Francisco J.; Dominguez Horna, Carlos (Consejo Superior de Investigaciones Cientificas, Spain). Span. ES 2146128 A1 20000716, 9 pp. (Spanish). CODEN: SPXXAD. APPLICATION: ES 1995-645 19950331. AB The process consists of adhesion of a photoresin to a substrate, such as Si oxide, using an organosilane, e.g., trimethoxy Pr silyl methacrylate (MPTS) or 3-aminopropyl triethoxysilane, silylation by submersion of the wafers of Si in a soln. of MPTS in toluene combined with .gtoreg.0.5 vol.% water, holding at <90.degree. for approx. 4 h, rinsing with MEK, and depositing the polyimide layer by centrifuging; photopolymn. of the organosilane to form a polysiloxane resulting from contact with moisture and crosslinking by photochem. excitation with UV light; producing pos. and neg. images depending on the order of the sequence of steps; and development by reactive ion etching using an O plasma after imidation of the polymer by heating at <400.degree.. The process gives pos. or neg. patterns with respect to the mask using the same photosensitive precursors and depending only on the order of exposure and silylation. It can be used for photodefinition of polymer layers of any thickness and applied to encapsulation of chem. sensors, multichip modules, microelectronic technol., electronic sensors, and integrated optics.

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TΤ
     919-30-2, 3-Aminopropyltriethoxysilane
        (substrate silylation agent; photomasking of photosensitive
        polyimides with organometallic compds. for photolithog.
        with oxygen plasma reactive ion etching)
RN
     919-30-2 HCAPLUS
     1-Propanamine, 3-(triethoxysilyl)- (9CI) (CA INDEX NAME)
CN
     OEt
Eto-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>
     OEt
IC
     ICM
          H01L021-312
          G03F007-037; G03F007-038; G03F007-075; G03F007-029; G03F007-36;
     ICS
          C08L079-08; C08K005-3417; C08K005-54
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and
CC
     Other Reprographic Processes)
     Section cross-reference(s): 76
     organosilane silylation photoresin adhesion photomasking; oxygen
ST
     plasma reactive ion etching photolithog
IT'
     Sputtering
        (etching, reactive, oxygen; photomasking of photosensitive
        polyimides with organometallic compds. for photolithog.
        with oxygen plasma reactive ion etching)
     Silanes
IT
        (organosilanes, substrate silylation agent; photomasking of
        photosensitive polyimides with organometallic compds. for
        photolithog. with oxygen plasma reactive ion etching)
     Photolithography
IT
       Photoresists
     Silvlation
        (photomasking of photosensitive polyimides with organometallic
        compds. for photolithog. with oxygen plasma reactive
        ion etching)
IT
     Polyimides, processes
     Polysiloxanes, processes
        (photomasking of photosensitive polyimides with organometallic
        compds. for photolithog. with oxygen plasma reactive
        ion etching)
IT
     Etching
        (sputter, reactive, oxygen; photomasking of photosensitive
        polyimides with organometallic compds. for photolithog.
        with oxygen plasma reactive ion etching)
     7782-44-7, Oxygen, uses
IT
        (plasma; photomasking of photosensitive polyimides with
        organometallic compds. for photolithog. with oxygen
        plasma reactive ion etching)
     919-30-2, 3-Aminopropyltriethoxysilane 2530-85-0
IT
        (substrate silylation agent; photomasking of photosensitive
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polyimides with organometallic compds. for **photolithog**. with oxygen plasma reactive ion etching)

L50 ANSWER 2 OF 11 HCAPLUS COPYRIGHT 2002 ACS
2001:293627 Document No. 134:289035 Method of patterning
polyamic acid layers. Beilin, Solomon I.; Chou, William T.; Ngo,
David Dung (Fujitsu Limited, Japan). U.S. US 6221567 B1 20010424, 9
pp. (English). CODEN: USXXAM. APPLICATION: US 1999-226974
19990108. PRIORITY: US 1998-PV71417 19980114.

Methods of etching polyamic acid layers and the like are disclosed. AB In exemplary embodiments of the present invention, the polymeric acid layer to be etched is alternatively exposed to etchant solns. (etchants) and rinse solns., where the etchant solns. are of relatively moderate alky. and the rinse solns. have a lower pH than the etchant solns. The present invention enables polymeric acid layers to be developed with std. basic etchants at relatively moderate concns. and at room temp. with little, if any, corrosion to any underlying metal layers. The present invention enables the more reliable and cleaner spin-spray processing method to be employed, thereby significantly increasing yields and reducing overall processing costs. The present invention also enables the etching of thick layers of polymeric acid without the need for special treatments, such as exposure to highly concd. etchant solns. or high temp. processing conditions.

IT 919-30-2, VM 651

(adhesion promoter; method of **patterning** polyamic acid layers)

RN 919-30-2 HCAPLUS

CN 1-Propanamine, 3-(triethoxysilyl)- (9CI) (CA INDEX NAME)

IC ICM G03F007-00

NCL 430323000

CC 76-3 (Electric Phenomena)

ST photolithog polyamic acid film etching

IT Etching

Photolithography

(method of **patterning** polyamic acid layers)

IT Polyamic acids

(method of patterning polyamic acid layers)

IT 919-30-2, VM 651

(adhesion promoter; method of **patterning** polyamic acid layers)

IT 7440-21-3, Silicon, uses 7631-86-9, Silica, uses (method of **patterning** polyamic acid layers)

IT 29319-22-0, PI2611

(polyamic acid coating; method of **patterning** polyamic acid layers)

IT 97396-58-2, AZ4620

(pos. photoresist; method of patterning polyamic acid layers)

L50 ANSWER 3 OF 11 HCAPLUS COPYRIGHT 2002 ACS

- 1999:101258 Document No. 130:149546 Novel methods of attaching probes to a solid support and uses thereof. Okamoto, Tadashi; Yamamoto, Nobuko; Suzuki, Tomohiro (Canon Kabushiki Kaisha, Japan). Eur. Pat. Appl. EP 895082 A2 19990203, 43 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO. (English). CODEN: EPXXDW. APPLICATION: EP 1998-306107 19980731. PRIORITY: JP 1997-207837 19970801; JP 1997-287046 19971020: JP 1998-209923 19980724.
- AB Provided is a method of attaching probes to a solid support in a markedly high d. and efficiency. An extremely small amt. of probe is contained within a liq., and droplets of the liq. are delivered to the solid support via an ink jet ejection method, thereby forming spots which contain the probe. Since one or more substances can bind specifically to target probes and said probes are arranged in a large no. on a solid support, the method can be used to swiftly and accurately det. a base sequence of a nucleic acid or detect a target nucleic acid in a sample.

1760-24-3, KBM603 2530-83-8, KBM403

(novel methods of attaching probes to a solid support and uses thereof)

RN 1760-24-3 HCAPLUS

CN 1,2-Ethanediamine, N-[3-(trimethoxysilyl)propyl]- (9CI) (CA INDEX NAME)

OMe 
$$\begin{array}{c} \text{OMe} \\ \mid \\ \text{MeO-Si-} (\text{CH}_2)_3 - \text{NH-CH}_2 - \text{CH}_2 - \text{NH}_2 \\ \mid \\ \text{OMe} \end{array}$$

RN 2530-83-8 HCAPLUS

CN Silane, trimethoxy[3-(oxiranylmethoxy)propyl]- (9CI) (CA INDEX NAME)

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IC
     ICM G01N033-543
     ICS
         C12Q001-68; C07K017-14; C12N011-14; G01N033-53; G01N033-68
CC
     3-1 (Biochemical Genetics)
     Section cross-reference(s): 9
     Photoresists
IT
        (UV; novel methods of attaching probes to a solid support and
        uses thereof)
IT
     Phenolic resins, analysis
        (arom. azide resist; novel methods of attaching probes
        to a solid support and uses thereof)
     Phenolic resins, analysis
IT
        (novolak, diazonaphthoquinone resist; novel methods of
        attaching probes to a solid support and uses thereof)
IT
     Polysulfones, analysis
        (poly(methylene sulfone); radiation decompn. resist;
        novel methods of attaching probes to a solid support and uses
        thereof)
IT
     Isoprenoids
        (polyisoprenoids, cyclized, arom. bisazide resist;
        novel methods of attaching probes to a solid support and uses
     Photolithography
IT
        (use in formation of matrix pattern; novel methods of
        attaching probes to a solid support and uses thereof)
     59269-51-1, Polyvinylphenol
                                   75742-13-1
IT
        (DEEP-UV resist contg.; novel methods of attaching
        probes to a solid support and uses thereof)
     25067-05-4, Polyglycidyl methacrylate
                                              203526-64-1, BK 739P
IT
        (DEEP-UV resist; novel methods of attaching probes to a
        solid support and uses thereof)
     80500-54-5, 2-Nitrobenzyl cholate
IT
        (dissoln. suppressant resist; novel methods of
        attaching probes to a solid support and uses thereof)
     1760-24-3, KBM603 2530-83-8, KBM403
TT
        (novel methods of attaching probes to a solid support and uses
        thereof)
     9011-14-7, Polymethyl methacrylate
IT
                                           25988-32-3,
     Polymethylisopropenyl ketone 76643-44-2, Polyhexafluorobutyl
     methacrylate 87842-32-8D, brominated
        (radiation decompn. resist; novel methods of attaching
        probes to a solid support and uses thereof)
     ANSWER 4 OF 11 HCAPLUS COPYRIGHT 2002 ACS
L50
              Document No. 130:8817 Fabrication of colloidal gold
     micropatterns using photolithographed self-assembled
     monolayers as templates. Liu, Jun-Fu; Zhang, Lin-Gang; Gu, Ning; Ren, Ji-Yun; Wu, Yan-Peng; Lu, Zu-Hong; Mao, Pan-Song; Chen, De-Ying
     (National Laboratory of Molecular and Biomolecular Electronics,
     Southeast University, Nanjing, 210096, Peop. Rep. China).
     Solid Films, 327-329, 176-179 (English) 1998. CODEN: THSFAP.
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0040-6090. Publisher: Elsevier Science S.A..

This paper provides a new method for the patterned AB metalization of self-assembled monolayers. Colloidal Au micropatterns were formed on photolithographed self-assembled monolayers (SAMs) of (3-mercaptopropyl)trimethoxysilane (MTS) on SiO2-coated Si substrates. As confirmed by SEM, the colloidal Au closely replicated the mask features. force microscopy (AFM) showed that the colloidal Au particles on the unirradiated monolayers of (3-mercaptopropyl)-trimethoxysilane were connected and an almost continuous monolayer of Au nanoparticles was X-ray photoelectron microscopy (XPM) revealed that the -SH terminal groups of the MTS monolayer in the exposed region were oxidized and the self-assembly of colloidal Au onto the unexposed region were through Au-S covalent bounding. The present approach may be applicable to assembly of microelectronic circuits and microbiosensors.

1T 4420-74-0, (3-Mercaptopropyl)-trimethoxysilane (fabrication of colloidal gold micropatterns using photolithographed self-assembled monolayers of (mercaptopropyl)trimethoxysilane on SiO2-coated Si substrates as templates)

RN 4420-74-0 HCAPLUS

CN 1-Propanethiol, 3-(trimethoxysilyl)- (7CI, 8CI, 9CI) (CA INDEX NAME)

$$^{
m OMe}$$
  $^{\mid}$   $^{\mid}$  MeO-Si-(CH<sub>2</sub>)<sub>3</sub>-SH  $^{\mid}$  OMe

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST colloidal gold micropattern **photolithographed** self assembled silane monolayer

IT Bond

(covalent; metalization of self-assembled monolayers of mercaptopropyltrimethoxysilane through covalent bonding of Au colloids to these **photolithographed** layers)

IT Nanoparticles
Photolithography

Self-assembled monolayers

(fabrication of colloidal gold micropatterns using photolithographed self-assembled monolayers of

(mercaptopropyl)trimethoxysilane on SiO2-coated Si substrates as templates)

IT Surface reaction

(metalization of self-assembled monolayers of mercaptopropyltrimethoxysilane through covalent bonding of Au colloids to these **photolithographed** layers)

IT Coating process

(metalization; metalization of self-assembled monolayers of mercaptopropyltrimethoxysilane through covalent bonding of Au colloids to these **photolithographed** layers)

IT Biosensors

(microbiosensors; fabrication of colloidal gold micropatterns using **photolithographed** self-assembled monolayers of (mercaptopropyl)trimethoxysilane on SiO2-coated Si substrates as templates in relation to)

IT Electric circuits

(microcircuits; fabrication of colloidal gold micropatterns using **photolithographed** self-assembled monolayers of (mercaptopropyl)trimethoxysilane on SiO2-coated Si substrates as templates in relation to)

IT 7440-57-5D, Gold, surface bonded, processes

(colloidal; fabrication of colloidal gold micropatterns using **photolithographed** self-assembled monolayers of (mercaptopropyl)trimethoxysilane on SiO2-coated Si substrates as templates)

IT 4420-74-0, (3-Mercaptopropyl)-trimethoxysilane 7440-21-3,

Silicon, processes

(fabrication of colloidal gold micropatterns using photolithographed self-assembled monolayers of (mercaptopropyl)trimethoxysilane on SiO2-coated Si substrates as templates)

- L50 ANSWER 5 OF 11 HCAPLUS COPYRIGHT 2002 ACS
- 1997:609956 Document No. 127:285940 Thermosetting electrodeposition coating composition having tackiness for transfer printing and pattern formation method using the same. Doi, Yorio; Nishida, Reijiro; Kawai, Kenzo Ro (Kansai Paint Co., Ltd., Japan; Dainippon Printing Co., Ltd.). Jpn. Kokai Tokkyo Koho JP 09235493 A2 19970909 Heisei, 17 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1996-65220 19960228.
- AB The coating compn. contains (A) a resin with OH and an ionic group and (B) a hardener with av. .gtoreq.2 epoxy-contg. functional groups having a cycloaliph. and/or bridged cycloaliph. backbone. The process consists of (1) forming an electrode with a pattern of an elec. insulating film on a conductive side of a substrate, (2) forming a releasing layer if necessary, (3) plating a metal, (4) soaking the substrate in the coating compn. to form an electrodeposition film, and (5) adhering a substrate on the coating layer, releasing, and transfering the coating and plating layers. The transfered coating layer is suitable for an etching resist mask.
- 156637-69-3P, (.gamma.-Glycidoxypropyl)trimethoxysilanemethyltrimethoxysilane copolymer

(releasing layer; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

RN 156637-69-3 HCAPLUS

CN Silane, trimethoxymethyl-, polymer with trimethoxy[3-(oxiranylmethoxy)propyl]silane (9CI) (CA INDEX NAME) CM 1

CRN 2530-83-8 CMF C9 H20 O5 Si

$$CH_2-O-(CH_2)_3-Si-OMe$$
OMe
OMe
OMe
OMe

CM 2

CRN 1185-55-3 CMF C4 H12 O3 Si

IC ICM C09D005-44 ICS C25D013-06

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST electrodeposition coating thermosetting resin tackiness photolithog; photoresist electrodeposition coating pattern transfer; polyepoxide hardener electrodeposition coating photolithog; hydroxy ionic resin thermosetting electrodeposition coating; vinylcyclohexane diepoxide polymer electrodeposition coating

IT Epoxy resins, uses

(alicyclic, hardeners; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

IT Epoxy resins, preparation

(cycloaliph. epoxide-crosslinked; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

IT Crosslinking agents

(cycloaliph. epoxides; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

IT Coatings

Electrodeposition

Photolithography

Photoresists

(epoxy resin electrodeposition coating compn. having tackiness

for photolithog.)

IT Polysiloxanes, preparation

(releasing layer; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

IT 196810-60-3P, Bisphenol A-bisphenol A diglycidyl ether-EHPE 3150-(3,4-epoxycyclohexyl)methyl methacrylate copolymer diethanolamine salt

(crosslinked; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

IT 196810-57-8P, Bisphenol A-bisphenol A diglycidyl ether-EHPE 3150 copolymer diethanolamine salt

(cycloaliph. epoxide-crosslinked; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

IT 128703-08-2P, (3,4-Epoxycyclohexyl) methyl methacrylate homopolymer 196810-62-5P, 3,4-Epoxytetrahydrobenzyl alcohol-HMDI isocyanurate copolymer 196810-65-8P

(hardeners; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

IT 25086-25-3, EHPE 3150

(hardeners; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)

- 156637-69-3P, (.gamma.-Glycidoxypropyl)trimethoxysilanemethyltrimethoxysilane copolymer 196810-67-0P, (.gamma.-Hydroxypropyl)trimethoxysilane-maleic anhydride-(.gamma.methacryloxypropyl)trimethoxysilane-methyltrimethoxysilane copolymer (releasing layer; epoxy resin electrodeposition coating compn. having tackiness for **photolithog**.)
- L50 ANSWER 6 OF 11 HCAPLUS COPYRIGHT 2002 ACS
- 1997:299472 Document No. 126:285347 High-resistance black radiation-sensitive resin composition, black hardened film, and black image formation. Koyanagi, Takao; Kato, Yoshinori; Imaizumi, Naoko; Kitaori, Tomoyuki; Futamura, Nobuyuki; Fukunaga, Masanori (Nippon Kayaku Kk, Japan). Jpn. Kokai Tokkyo Koho JP 09054431 A2 19970225 Heisei, 9 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1995-232126 19950818.
- The title resin compn. contains a radiation-sensitive resin contg. a AB radiation-crosslinkable compd. and/or a resin and an optional other resin, Ti black surface treated with a silane compd., and optionally a photopolymn. initiator and an org. pigment obtained by mixing carbon black with other colorant to make black. A black hardened film obtained by using the compn. and a black image-forming method by photolithog. using the compn. are The compn. useful in prepn. of a black matrix shows also claimed. high resistance, photosensitivity, light-shielding properties, and dispersion stability. Thus, a radiation-sensitive resin compn. was prepd. by mixing .gamma.-glycidoxypropyltrimethoxysilane-treated Ti black with a compn. contg. DPHA (radiation-crosslinkable resin), Me methacrylate-methacrylic acid-benzyl methacrylate-2-hydroxyethyl methacrylate copolymer, and photopolymn. initiators. 2530-83-8 IT

(high-resistance radiation-sensitive resin compn. for black image

formation)

RN 2530-83-8 HCAPLUS

Silane, trimethoxy[3-(oxiranylmethoxy)propyl]- (9CI) (CA INDEX CN

$$CH_2-O-(CH_2)_3-Si-OMe$$
OMe
OMe
OMe
OMe
OMe
OMe

IC ICM G03F007-004

> C08K009-06; C08L101-00; C09D005-00; C09D007-12; C09D201-00; ICS G02B005-20; G02F001-1335; G03F007-027; H01J009-227

74-5 (Radiation Chemistry, Photochemistry, and Photographic and CC Other Reprographic Processes)

Section cross-reference(s): 37

radiation sensitive black resin high resistance; resist ST radiation sensitive black image

Resists IT

> (radiation-sensitive; high-resistance radiation-sensitive resin compn. for black image formation)
> 2530-83-8
> 3388-04-3. heta -(3.4-

TT

Epoxycyclohexyl)ethyltrimethoxysilane 144113-99-5, Titanium Black 13R

(high-resistance radiation-sensitive resin compn. for black image formation)

ANSWER 7 OF 11 HCAPLUS COPYRIGHT 2002 ACS L50

Document No. 125:99862 Planar compatible polymer 1996:376423 Munoz, J.; technology for packaging of chemical microsensors. Bratov, A.; Mas, R.; Abramova, N.; Dominguez, C.; Bartroli, J. (Centro Nacional Microelectronica-CSIC, Bellaterra, 08193, Spain). J. Electrochem. Soc., 143(6), 2020-2025 (English) 1996. CODEN: JESOAN. ISSN: 0013-4651.

By applying special thick film photolithog. it is possible AB to realize a highly automative ion-sensitive field effect transistor (ISFET) packaging at the wafer level. In this paper two approaches based on photolithog processing of

encapsulating layers are presented. A lift-off method of com. thermo-curable encapsulants has been investigated, as well as direct photopolymn. of UV-curable encapsulant compns. contg. either epoxy acrylate or polyurethane acrylate oligomers together with monomers and photoinitiators. Lift-off has been developed using either thin or thick photoresist sacrificial layers. The best compatibility implies thick photosensitive polyimide layers as a sacrificial photoresist together with the encapsulant

material based on alumina-filled epoxy. However, better results are obtained by using photopatternable encapsulant polymers that permit

application of thin or thick polymer layers on a wafer substrate contg. ISFET chips. Windows over the gate region and contact pads are opened by exposure to UV light in a std. mask aligner system. Compns. based on epoxy acrylate oligomers are proved to be more reliable in a packaging process. Lifetime of encapsulated micro-sensors for pH measurements is presented.

919-30-2, 3-Triethoxysilyl-1-propylamine
(silylating agent; highly automative ion-sensitive field effect transistor packaging at wafer level by special thick film photolithog)

RN 919-30-2 HCAPLUS

CN 1-Propanamine, 3-(triethoxysilyl)- (9CI) (CA INDEX NAME)

OEt 
$$|$$
 EtO-Si-(CH<sub>2</sub>)<sub>3</sub>-NH<sub>2</sub>  $|$  OEt

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 36, 76

insulating ISFET packaging microelectronics sacrificial photoresist; thick film photolithog chem microsensor packaging; encapsulated microsensor pH ISFET insulating packaging

IT Polyimides, uses

(thick photosensitive polyimide layers as sacrificial photoresist in ISFET packaging at wafer level)

IT Transistors

(field-effect, ion-selective, highly automative ion-sensitive field effect transistor packaging at wafer level by special thick film **photolithog**.)

IT Lithography

(**photo**-, highly automative ion-sensitive field effect transistor packaging at wafer level by special thick film **photolithog**.)

IT Resists

(photo-, photoresist sacrificial layers in ISFET packaging at wafer level by special thick film photolithog.)

IT 118731-53-6, Probimide 348 126904-05-0, Selectilux HTR 3-200 153191-91-4, Probimide 408

(resist. photoresist sacrificial layers in

(resist; photoresist sacrificial layers in
ISFET packaging at wafer level by special thick film
photolithog;)

- 919-30-2, 3-Triethoxysilyl-1-propylamine 2530-85-0
   (silylating agent; highly automative ion-sensitive field effect
   transistor packaging at wafer level by special thick film
   photolithog.)
- L50 ANSWER 8 OF 11 HCAPLUS COPYRIGHT 2002 ACS
- 1995:1002097 Document No. 124:101674 A top-surface imaging approach based on the light-induced formation of dry-etch barriers. Schaedeli, U.; Hofmann, M.; Tinguely, E.; Muenzel, N. (Materials Research, Ciba-Geigy Inc., Marly/Fribourg, 1723, Switz.). ACS Symp. Ser., 614 (Microelectronics Technology), 299-317 (English) 1995. CODEN: ACSMC8. ISSN: 0097-6156.
- It is believed that optical lithog. will continue to play an AB important role for the manufg. of future generation IC devices. Surface imaging processes might be required to overcome the small depth of focus assocd. with new high numerical aperture exposure tools. Our new concept is based on the light induced formation of sites for the fixing of reactive monomers in the top zones of a highly absorbing resist film. These sites, which typically consist of photolytically generated Broensted acid or radicals, are able to oligomerize silicon-contg., reactive monomers during a baking step. The net result is the formation of a silicon gradient in the top zone of the exposed resist film, which subsequently acts as a dry etch barrier. Neg. tone images were obtained by exposing resist films, mainly composed from a linear matrix polymer, a sulfonium salt type photoacid generator, and silicon-contg. epoxy monomers, with light of 193 nm or 254 nm wavelength. **7422-52-8**

(top-surface imaging of **photoresists** based on photooligomerization of silicon contg. monomers)

RN 7422-52-8 HCAPLUS

CN Trisiloxane, 1,1,1,3,5,5,5-heptamethyl-3-[3-(oxiranylmethoxy)propyl](9CI) (CA INDEX NAME)

$$\begin{array}{c|c} O & O-SiMe_3\\ & | \\ CH_2-O-(CH_2)_3-Si-Me\\ & | \\ O-SiMe_3 \end{array}$$

- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- ST surface imaging photoresist photooligomerization silicon

```
monomer
IT
     Etching
        (top-surface imaging of photoresists based on
        photooligomerization of silicon contq. monomers)
IT
     Polymerization
        (oligomerization, photochem., top-surface imaging of
        photoresists based on photooligomerization of silicon
        contg. monomers)
     Lithography
IT
       Resists
        (photo-, top-surface imaging of photoresists
        based on photooligomerization of silicon contg. monomers)
TT
     2362-10-9P
                  76104-14-8P
                                90492-23-2P
                                              101667-50-9P
     172667-45-7P
        (top-surface imaging of photoresists based on
        photooligomerization of silicon contg. monomers)
IT
     172667-46-8P
        (top-surface imaging of photoresists based on
        photooligomerization of silicon contg. monomers)
IT
     126-80-7, 1,3-Bis(3-glycidyloxypropyl)tetramethyldisiloxane
     4515-14-4 7422-52-8
                           18415-68-4
                                        65842-29-7
                                                      90492-24-3
                                 162546-55-6
     162546-50-1
                   162546-52-3
                                                162546-56-7
        (top-surface imaging of photoresists based on
        photooligomerization of silicon contg. monomers)
     ANSWER 9 OF 11 HCAPLUS COPYRIGHT 2002 ACS
L50
              Document No. 123:270507 New top surface imaging approach
1995:672250
     based on the light induced built-up of dry etch barriers.
     Schaedeli, U.; Hofmann, M.; Tinguely, E.; Munzel, N. (Materials
     Research, Ciba-Geigy Corp., Fribourg, 1723, Switz.).
                                                            Polym. Mater.
     Sci. Eng., 72, 106-7 (English) 1995. CODEN: PMSEDG.
     0743-0515.
     A top surface imaging approach, which yields neg. tone images of the
AB
     mask, is presented. It is based on the light induced generation of
     mol. filters in the top zones of the resist film, which
     are able to retard silicon contg. reactive monomers by rapid
     oligomerization. The resulting dry etch barriers allow for a
     pattern transfer down to the substrate by means of oxygen
     plasma etching. 7422-52-8
IT
        (top surface imaging approach which yields neg. tone images of
        mask based on light induced built-up of dry etch barriers)
RN
     7422-52-8 HCAPLUS
     Trisiloxane, 1,1,1,3,5,5,5-heptamethyl-3-[3-(oxiranylmethoxy)propyl]-
CN
      (9CI) (CA INDEX NAME)
```

$$\begin{array}{c|c} \text{O-SiMe}_3\\ & |\\ \text{CH}_2\text{-O-(CH}_2)_3\text{-Si-Me}\\ & |\\ \text{O-SiMe}_3\\ \end{array}$$

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 76

ST top surface imaging photomask etching photolithog

IT Lithography

Resists

(**photo**-, top surface imaging approach which yields neg. tone images of mask based on light induced built-up of dry etch barriers)

IT 126-80-7, 1,3-Bis(3-glycidyloxypropyl)tetramethyldisiloxane 4515-14-4 **7422-52-8** 18415-68-4 162546-52-3 162546-55-6 162546-56-7

(top surface imaging approach which yields neg. tone images of mask based on light induced built-up of dry etch barriers)

L50 ANSWER 10 OF 11 HCAPLUS COPYRIGHT 2002 ACS

1993:69701 Document No. 118:69701 Manufacture of color filters for liq.-crystal display devices. Iwamoto, Masao; Kimura, Kuniko (Toray Industries, Inc., Japan). Jpn. Kokai Tokkyo Koho JP 04151604 A2 19920525 Heisei, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1990-277249 19901015.

The manufg. process comprises the steps of: forming a laminate contg. (1) a transparent substrate, (2) a patterned resist, (3) a dyeable, (4) a photosensitive, and (5) a Si-rubber layer; forming the patterned apertures by selectively etching (3)/(4) photolithog.; forming the red/green/blue pixel elements by dying (2) exposed in the apertures; and removing the patterned (3)/(4). The filter has an improved surface integrity and a superior color rendering.

IT 919-30-2, A 1100

(color filters from, for liq.-crystal displays, manuf. of)

RN 919-30-2 HCAPLUS

CN 1-Propanamine, 3-(triethoxysilyl)- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} \text{OEt} \\ \vdots \\ \text{EtO-Si-} (\text{CH}_2)_3 - \text{NH}_2 \\ \vdots \\ \text{OEt} \end{array}$$

IC ICM G02B005-20

ICS G09F009-30 CC 73-12 (Optical, Electron, and Mass Spectroscopy and Other Related Properties) 919-30-2, A 1100 75882-39-2, Sumiresin PR 50235 IT (color filters from, for lig.-crystal displays, manuf. of) ANSWER 11 OF 11 HCAPLUS COPYRIGHT 2002 ACS L50 Photolithography on Document No. 117:201695 1992:601695 microstructured surfaces using photosensitive polyimide. Munoz, J.; Van Hal, R. E. G.; Bergveld, P. (MESA Res. Inst., Univ. Twente, Enschede, 7500 AE, Neth.). Sens. Mater., 3(6), 345-58 (English) CODEN: SENMER. ISSN: 0914-4935. Improvements in patterning microstructured surfaces by AB photosensitive polyimides as an alternative to std. pos. and neg. photoresists are presented. Two different types of polyimide photoresists are used to check step coverage and image transfer to specific test substrates. These test substrates contain mesalike steps, which can be seen as a general configuration of 3-dimensional Si sensors. Step coverage is characterized by leakage-current measurements and scanning electron microscope (SEM) spectroscopy. Thin photosensitive polyimide layers showed better step coverage and resoln. than std. photoresist Thick photosensitive polyimide showed good planarization properties when 2 layers were used. However, it is necessary to use a combination of wet and dry etching to obtain good resoln. 919-30-2, 3-Aminopropyl triethoxy silane IT (silylating agent, for photosensitive polyimide in photolithog ) RN919-30-2 HCAPLUS 1-Propanamine, 3-(triethoxysilyl)- (9CI) CN(CA INDEX NAME) OEt Eto-Si- $(CH_2)_3$ -NH<sub>2</sub> OEt . 74-5 (Radiation Chemistry, Photochemistry, and Photographic and CC

Other Reprographic Processes) IT Polyimides, uses (photosensitive, photolithog. on microstructured surfaces using) Lithography IT (**photo**, on microstructured surfaces using polyimide) Resists TT (photo-, photosensitive polyimides as, photolithog. on microstructured surfaces using) 9077-64-9, Waycoat IC 97445-31-3, S 1400-31. IT (lithog. patterning of microstructured surfaces by, photosensitive polyimides as alternative to)

- IT 126904-05-0, Selectilux HTR 3-200 135153-00-3, Probimide 412 (photolithog. on microstructured surfaces using)
- IT 999-97-3, Hexamethyl disilazane (silylating agent, for neg. photoresist in photolithog.)
- 919-30-2, 3-Aminopropyl triethoxy silane (silylating agent, for photosensitive polyimide in photolithog.)

CI COM

LC STN Files: AGRICOLA, ANABSTR, BEILSTEIN\*, BIOBUSINESS, BIOSIS,
BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CHEMCATS,
CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, CSNB, DETHERM\*, EMBASE, HODOC\*,
HSDB\*, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MSDS-OHS, NIOSHTIC, PIRA,
PROMT, RTECS\*, SPECINFO, TOXCENTER, USPAT2, USPATFULL
(\*File contains numerically searchable property data)
Other Sources: DSL\*\*, EINECS\*\*, TSCA\*\*
(\*\*Enter CHEMLIST File for up-to-date regulatory information)

=>

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

7755 REFERENCES IN FILE CA (1962 TO DATE)
1179 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
7772 REFERENCES IN FILE CAPLUS (1962 TO DATE)
92 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

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ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS
L3
     919-30-2 REGISTRY
RN
     1-Propanamine, 3-(triethoxysilyl)- (9CI) (CA INDEX NAME)
CN
OTHER CA INDEX NAMES:
     Propylamine, 3-(triethoxysilyl)- (6CI, 8CI)
OTHER NAMES:
     (.gamma.-Aminopropyl)triethoxysilane
CN
CN
     .gamma.-Triethoxysilylpropylamine
     3-(Triethoxysilyl)-1-propanamine
CN
     3-(Triethoxysilyl)-1-propylamine
CN
     3-(Triethoxysilyl)propanamine
CN
     3-(Triethoxysilyl)propylamine
CN
     3-Aminopropyltriethoxysilane
     3-APTES
CN
CN
     A 0750
     A 1100
CN
CN
     A 1100 (coupling agent)
     A 1102
CN
CN
     A 1102 (silane derivative)
CN
     A 1112
CN
     A 1112 (coupling agent)
     AGM 3
CN
CN
     AGM 9
CN
     AMEO
CN
     Aminopropyltriethoxysilane
CN
     AP 1690
CN
     APS
     APS (coupling agent)
CN
CN
     APS-E
     APTES
CN
     DS-AMEO
CN
CN
     Dynasylan 1203
CN
     Dynasylan AMEO
CN
     Dynasylan AMEO-T
     GF 93
CN
CN
     Hydrosil 2627
CN
     KBE 903
     KH 550
CN
     NB 1114
CN
     NUCA 1100
CN
     Prosil 220
CN
CN
     Prosil 221
CN
     s 330
     Sila-Ace S 330
CN
     Silane AMG 9
CN
CN
     Silicone A 1100
CN
     Silquest A 1100
CN
     Silquest A 1102
CN
     Triethoxy(.gamma.-aminopropyl)silane
CN
     Triethoxy(3-aminopropyl)silane
CN
     TSC 202
CN
     TSL 8331
CN
     Unisilan 13
     VM 651
     VM 651 (coupling agent)
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
FS
     3D CONCORD
     12738-50-0, 131641-77-5, 60000-97-7, 96726-79-3, 106096-79-1, 71618-18-3,
DR
     143178-71-6, 86836-28-4, 159778-17-3, 204987-58-6
MF
     C9 H23 N O3 Si
```